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6	19	29.2	20434	2	AC141151	Rattus norvegicus
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ALIGNMENTS

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REFERENCE JOURNAL TITLE Direct Submission
Submitted (22-MAY-2002) Genome Sequencing Center, 4444 Forest Park
Parkway, St. Louis, MO 63108, USA
4 (bases 1 to 193659)
AUTHORS McPherson,J.D. and Waterston,R.H.
JOURNAL TITLE Direct Submission
Submitted (17-AUG-2002) Genome Sequencing Center, 4444 Forest Park
Parkway, St. Louis, MO 63108, USA
5 (bases 1 to 193659)
REFERENCE AUTHORS Wilson,R.
JOURNAL TITLE Direct Submission
Submitted (05-NOV-2003) Department of Genetics, Washington
University, 4444 Forest Park Avenue, St. Louis, Missouri 63108, USA
On Aug 17, 2002 this sequence version replaced gi:21105859.
COMMENT -----
Center: Washington University Genome Sequencing Center
Center code: WUGSC
Web site: http://genome.wustl.edu
Contact: submissions@waston.wustl.edu
----- Summary Statistics -----
Center project name: M_BB0216M06
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NOTICE: This sequence may not represent the entire insert of this
clone. It may be shorter because we only sequence overlapping
clone sections once, or longer because we provide a small overlap
between neighboring data submissions.

This sequence was finished as follows unless otherwise noted:
all regions were double stranded, sequenced with an alternate
chemistry, or covered by high quality data (i.e., phred quality >=
30); an attempt was made to resolve all sequencing problems, such
as compressions and repeats; all regions were covered by sequence
from more than one subclone; and the assembly was confirmed by
restriction digest.

MAPPING INFORMATION:
Mapping information for this clone was provided by Dr. Wes Warren,
Department of Genetics, Washington University, St Louis MO. For
additional information about the map position of this sequence, see
http://genome.wustl.edu

SOURCE INFORMATION:
The RPCI-24 BAC library has been constructed by Pieter de Jong and
coworkers (http://www.chori.org) from male C57BL/6J mouse spleen
and/or brain genomic DNA. The clone and detailed information can be
obtained from Pieter de Jong and coworkers at http://www.chori.org

NEIGHBORING SEQUENCE INFORMATION:
This sequence is the entire insert of the clone.
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Best Local Similarity	100.0%;	Pred. No. 4.8;		
Matches 19;	Conservative 0;	Mismatches 0;	Indels 0;	Gaps 0;

QY 40 AAACCTGCTGATGACAC 58
Db 95628 AAACCTGCTGATGACAC 95646

LOCUS	DEFINITION	AC141530	215983 bp	DNA	1 linear	HTG 17-MAR-2003
LOCUS	Rattus norvegicus clone CH230-517N11, WORKING DRAFT SEQUENCE, 47 unordered pieces.	AC141530	215983 bp	DNA	1 linear	HTG 17-MAR-2003

ACCESSION	AC141530.1
VERSION	AC141530.1 GI:28975794
KEYWORDS	HTG; HTGS_PHASE1; HTGS_DRAFT.
SOURCE	Rattus norvegicus (Norway rat)

ORGANISM	<i>Rattus norvegicus</i>
Eukaryota; Metazoa; Chordata; Craniata; Vertebrata; Euteleostomi;	
Mammalia; Eutheria; Rodentia; Sciurognathi; Muridae; Murinae;	
<i>Rattus</i> .	
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REFERENCE	Muzny, D., Marie, M., Metzker, M., Lee, A., Abramson, S., Adams, C., Alder, J.,
AUTHORS	

Allen, C., Allen, H., Albriocks, S., Amin, A., Anguano, D., Anyelebechi, V., Aoyagi, A., Ayodeji, M., Bacas, E., Baden, H., Baldwin, D., Banderamaika, D., Barber, M., Barnstead, M., Benhmed, F., Bialwo, K., Blair, J., Blankenburg, K., Blyth, P., Brown, M., Bryant, N., Buhay, C., Burch, C.P., Burrill, K., Calderon, E., Cardenas, V., Carter, K., Cavazos, I., Cessari, H., Center, A., Chacko, T., Chaver, D., Chen, G., Chen, R., Chen, Y., Chen, Z., Chu, J., Cleveland, C., Cockrell, R., Cox, C., Coyle, M., Crete, A., D'Souza, L., Davila, M.L., Davis, C., Davy-Carroll, L., De Anda, C., Dederich, D., Delgado, O., Benson, C., Delano, C., Ding, Y., Dinh, H., Dwyer, K., Draper, H., Dugan-Rocha, S., Dunn, A., Durdin, K., Duval, B., Evans, K., Egan, A., Escotto, M., Eugene, C., Evans, C.A., Falls, T., Fan, G., Fernandez, S., Finley, M., Flagg, N., Forbes, L., Foster, M., Foster, P.,

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 Holt, R.A., Smith, H.O., Weinstock, G. and Gibbs, R.A.

Unpublished
 2 (bases 1 to 215983)

Worley, K.C.

Direct Submission
 Submitted (17-MAR-2003) Human Genome Sequencing Center, Department
 of Molecular and Human Genetics, Baylor College of Medicine, One
 Baylor Plaza, Houston, TX 77030, USA

----- Genome Center
 Center: Baylor College of Medicine
 Center code: BCM
 Web site: <http://www.hgsc.bcm.tmc.edu/>
 Contact: hgsc-help@bcm.tmc.edu
 ----- Project Information
 Center project name: KRPJ
 Center clone name: CH230-517N11
 ----- Summary Statistics
 Sequencing vector: Plasmid
 Chemistry: Dye-terminator Big Dye 100% of reads
 Assembly program: Phrap, version 0.990329
 Consensus quality: 200142 bases at least Q40
 Consensus quality: 204905 bases at least Q30
 Consensus quality: 208606 bases at least Q20
 Estimated insert size: 2045.6; sum-of-contigs estimation
 Quality coverage: 3x in Q20 bases; sum-of-contigs estimation

NOTE: Estimated insert size may differ from sequence length
 (see http://www.hgsc.bcm.tmc.edu/sgcs/genbank_draft_data.html).
 NOTE: This is a "working draft" sequence. It currently
 consists of 47 contigs. The true order of the pieces
 is not known and their order in this sequence record is
 arbitrary. Gaps between the contigs are represented as
 runs of N, but the exact sizes of the gaps are unknown.
 This record will be updated with the finished sequence
 as soon as it is available and the accession number will
 be preserved.

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 * 1368 1467: gap of unknown length
 * 1468 2579: contig of 1112 bp in length

2580 2679: gap of unknown length
 * 2680 4117: contig of 1438 bp in length
 * 4118 4217: gap of unknown length
 * 4218 5365: contig of 1148 bp in length
 * 5366 5465: gap of unknown length
 * 5466 6161: contig of 1351 bp in length
 * 6162 6916: gap of unknown length
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 * 8048 9570: contig of 1523 bp in length
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 * 33359 36264: contig of 2907 bp in length
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 * 76753 83329: contig of 3859 bp in length
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 * 116372 116471: contig of 7376 bp in length
 * 116472 121822: gap of unknown length
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 * 121824 121822: gap of unknown length
 * 121825 127516: contig of 5594 bp in length
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Accession	Contig	Length
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160563	contig of 7705 bp in length	168867
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187193	contig of 14554 bp in length	201446
201447	gap of unknown length	201546
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ORIGIN

[illegible]

RESULT	6
AC1141151	
LOCUS	AC1141151
DEFINITION	AC1141151. Rattus norvegicus clone CH230-114H6, WORKING DRAFT SEQUENCE, 67
ACCSSION	AC1141151
VERSION	AC1141151.2
KEYWORDS	HTG; HTGS_PHASE1; HTGS_DRAFT.
SOURCE	Rattus norvegicus (Norway rat)
ORGANISM	Rattus norvegicus

REFERENCE
AUTHORS

(Pages 1 to 220434)

Mizuy, D., Marie, Metzger, M. Lee, Abramson, S., Adams, C., Alder, J., Allen, C., Allen, H., Alstrooks, S., Amin, A., Anguiano, D., Anyalebsch, V., Aoyagi, A., Aydeli, M., Baca, S., Baden, H., Baldwin, D., Bandaranaike, D., Barber, M., Barnstead, M., Beaumond, F., Belsalo, K., Blair, J., Blankenburg, K., Blyth, P., Brown, M., Bryant, N., Buhay, C., Burch, P., Butrell, K., Calderon, E., Cardenas, V., Carter, K., Cavazos, T., Cessari, H., Centar, A., Chacko, J., Chavez, D., Chen, G., Chen, R., Chen, Y., Chen, Z., Cleveland, C., Cockrell, R., Cox, C., Coyle, M., Cree, A., D'Souza, L., Davila, M., Davis, C., Davy-Carrillo, L., De Anda, C., Dederich, D., Delgado, O., Denison, S., Deramo, C., Ding, Y., Dinh, H., Ditya, K., Draper, H., Dugan-Rocha, S., Dunn, A., Dublin, K., Duval, B., Eaves, K., Egan, A., Escotto, M., Eugene, S., Evans, C.A., Falls, T., Fan, G., Fernandez, S., Finley, M., Flagg, N., Forbes, L., Foster, M., Foster, P., Fraser, C.M., Gabriel, A., Gantla, R., Garcia, A., Garner, T., Garza, M., Gebregiorgis, E., Geer, K., Gill, R., Grady, M., Guerra, M., Guevara, M., Gunaratne, P., Haaland, W., Hamil, C., Hamilton, C., Hamilton, K., Harvey, Y., Havlak, P., Hawes, A., Henderson, N., Hernandez, J., Hernandez, R., Hines, S., Hladun, S.L., Hodgson, A., Hognes, M., Hollins, B., Howells, S., Huily, S., Hume, J., Idebri, D., Jackson, A., Jackson, B., Jacob, L., Jiang, H., Johnson, B., Johnson, R., Jolyvet, A., Karpethy, S., Kelly, S., Kelly, S., Khan, Z., King, L., Kovar, C., Kowis, C., Kraft, C.L., Lebow, H., Leven, J., Lewis, L., Li, Z., Liu, J., Liu, J., Liu, W., Liu, R., London, P., Longacre, S., Lopez, J., Lorensunewa, L., Louised, H., Losado, R.-J., Lu, X., Lu, X.,

TITLE Direct Submission
JOURNAL Unpublished
REFERENCE 2 (bases 1 to 220434)
AUTHORS Morley, K.C.
TITLE Direct Submission
JOURNAL Submitted (10-MAR-2003) Human Genome Sequencing Center, Department

REFERENCE 3 (bases 1 to 220434)
AUTHORS Worley, K.C.
TITLE Direct Submission
JOURNAL Submitted (27-MAR-2003) Human Genome Sequencing Center, Department of Molecular and Human Genetics, Baylor College of Medicine, One Baylor Plaza, Houston, TX 77030, USA
COMMENT On May 14, 2003 this sequence version replaced gi:28894506.

```

* Chemistry. Dye-terminator Big Dye 3.0001 of
* Assembly program: Phrap, version 0.9901329
* Consensus quality: 192518 bases at least Q40
* Consensus quality: 201921 bases at least Q30
* Consensus quality: 208432 bases at least Q20
* Estimated insert size: 198182; sum-of-confid-
* Quality coverage: 3x in Q20 bases; sum-of-co-
*
* NOTE: Estimated insert size may differ from size
* (see http://www.hgscc.bcm.tmc.edu/docs/Genbaas/)
* NOTE: This is a 'working draft' sequence. It
* consists of 67 contigs. The true order of the
* is not known and their order in this sequence
* arbitrary. Gaps between the contigs are repre-
* runs of 'N', but the exact sizes of the gaps are
* This record will be updated with the finished
* as soon as it is available and the accession n-
* be preserved.
*
* 1 1220: contig of 1220 bp in length
*
* 1221 1320: gap of unknown length
*
* 1321 2711: contig of 1391 bp in length
*
* 2712 2811: gap of unknown length
*
* 2812 4402: contig of 1491 bp in length
*
* 4403 4402: gap of unknown length
*
* 4403 5435: contig of 1033 bp in length

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* 5436 5535: gap of unknown length
* 5536 6682: contig of 1147 bp in length
* 6683 6782: gap of unknown length
* 6783 8122: contig of 1343 bp in length
* 8126 8225: gap of unknown length
* 8226 9654: contig of 1428 bp in length
* 9654 9754: gap of unknown length
* 9754 10893: contig of 1139 bp in length
* 10893 10993: gap of unknown length
* 10993 12420: contig of 1427 bp in length
* 12420 12519: gap of unknown length
* 12519 13792: contig of 1273 bp in length
* 13792 13892: gap of unknown length
* 13893 15227: contig of 1335 bp in length
* 15228 15327: gap of unknown length
* 15328 16824: contig of 1496 bp in length
* 16824 16923: gap of unknown length
* 16923 18594: contig of 1671 bp in length
* 18594 18694: gap of unknown length
* 18694 20057: contig of 1363 bp in length
* 20057 20157: gap of unknown length
* 20158 21205: contig of 1048 bp in length
* 21206 21305: gap of unknown length
* 21306 22447: contig of 1142 bp in length
* 22448 22548: gap of unknown length
* 22548 24159: contig of 1611 bp in length
* 24159 24258: gap of unknown length
* 24258 25928: contig of 1671 bp in length
* 25928 26028: gap of unknown length
* 26029 27224: contig of 1195 bp in length
* 27225 27325: gap of unknown length
* 27325 28721: contig of 1397 bp in length
* 28722 28822: gap of unknown length
* 28822 29840: contig of 1019 bp in length
* 29841 29941: gap of unknown length
* 29941 31550: contig of 1609 bp in length
* 31550 31650: gap of unknown length
* 31650 33453: contig of 1803 bp in length
* 33453 33552: gap of unknown length
* 33552 35257: contig of 1705 bp in length
* 35258 35358: gap of unknown length
* 35358 37459: contig of 2102 bp in length
* 37460 37560: gap of unknown length
* 37560 38811: contig of 1251 bp in length
* 38811 38911: gap of unknown length
* 38911 39980: contig of 1069 bp in length
* 39980 40079: gap of unknown length
* 40080 41615: contig of 1536 bp in length
* 41616 41715: gap of unknown length
* 41716 44028: contig of 2313 bp in length
* 44029 44128: gap of unknown length
* 44129 45437: contig of 1308 bp in length
* 45437 45537: gap of unknown length
* 45537 48041: contig of 2505 bp in length
* 48042 48141: gap of unknown length
* 48142 50554: contig of 2413 bp in length
* 50555 50654: gap of unknown length
* 50655 52760: contig of 2106 bp in length
* 52761 52861: gap of unknown length
* 52862 54582: gap of unknown length
* 54583 54682: gap of unknown length
* 54683 57043: contig of 2361 bp in length
* 57044 57143: gap of unknown length
* 57144 59095: contig of 1952 bp in length
* 59096 59195: gap of unknown length
* 59196 62500: contig of 3305 bp in length
* 62501 62600: gap of unknown length
* 62601 64620: contig of 2020 bp in length
* 64621 64720: gap of unknown length
* 64721 67094: contig of 2374 bp in length
* 67095 68971: gap of unknown length
* 68972 69071: contig of 1777 bp in length
* 69072 69171: gap of unknown length

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* 69072 71750: contig of 2679 bp in length
* 71751 71850: gap of unknown length
* 71851 74352: contig of 2502 bp in length
* 74353 74452: gap of unknown length
* 74453 77263: contig of 2811 bp in length
* 77264 77363: gap of unknown length
* 77364 80267: contig of 2903 bp in length
* 80268 80366: gap of unknown length
* 80367 83477: contig of 3111 bp in length
* 83478 83577: gap of unknown length
* 83578 86160: contig of 2583 bp in length
* 86161 86260: gap of unknown length
* 86261 88294: contig of 2034 bp in length
* 88295 88394: gap of unknown length
* 88395 93403: contig of 5009 bp in length
* 93404 93503: gap of unknown length
* 93504 98849: contig of 5346 bp in length

Query Match      29.2% Score 19; DB 2; Length 220434;
Best Local Similarity 100.0%; Pred. No. 4.7;
Matches 19; Conservative 0; Mismatches 0; Indels 0; Gaps 0;

QY      12 ATCTGCCGCTCTTCCAGC 30
Db      26915 ATCTGCCGCTCTTCCAGC 26933

RESULT 7
AL445257
LOCUS      Homo sapiens chromosome 1 clone RP5-1175N1, 14 unordered pieces.
DEFINITION
AL445257.1 GI:10716509
ACCESSION
AL445257.1
VERSION
HTG; HTGS PHASE1; HTGS_CANCELLED.
KEYWORDS
Homo sapiens (human)
SOURCE
Homo sapiens
Eukaryota; Metazoa; Chordata; Craniata; Vertebrata; Euteleostomi;
Mammalia; Eutheria; Primates; Catarrhini; Homnidae; Homo.
1
REFERENCE
  Burton, J.
  Direct Submission
  Submitted (12-JUN-2001) Sanger Centre, Hinxton, Cambridgeshire,
  CB10 1SA, UK. E-mail enquiries: humquery@sanger.ac.uk
  requester: clonerequest@sanger.ac.uk
  ----- Genome Center
  Center: Sanger Centre
  Center code: SC
  Web site: http://www.sanger.ac.uk
  Contact: humquery@sanger.ac.uk
  ----- Project Information
  Center project name: d11175N1
  ----- Summary Statistics
  Assembly program: XAPP4; Version 4.5
  Sequencing vector: plasmid; L08752; 100% of reads
  Chemistry: Dye-terminator Big Dye; 100% of reads
  Consensus quality: 28716 bases at least Q40
  Consensus quality: 32676 bases at least Q30
  Consensus quality: 35450 bases at least Q20
  Insert size: 38286; sum-of-contigs
  Insert size: 138118; 12.9% error; agarose-fp
  Quality coverage: 1.51x in Q20 bases; sum-of-contigs Quality
  coverage: 0.97x in Q20 bases; agarose-fp
  -----
  * NOTE: This is a 'working draft' sequence. It currently
  * consists of 14 contigs. The true order of the pieces
  * is not known and their order in this sequence record is
  * arbitrary. Gaps between the contigs are represented as
  * runs of N, but the exact sizes of the gaps are unknown.
  * This record will be updated with the finished sequence
  * as soon as it is available and the accession number will
  * be preserved.
  * 1
  * 2942: contig of 2942 bp in length
  * 2943 3042: gap of 100 bp

```

FEATURES	Location/Qualifiers
source	1. .39596

ORIGIN

LOCUS	AL35713	103513 bp	DNA	linear	PR1	01-MAR-2001
DEFINITION	Human DNA sequence from clone RP11-3J23 on chromosome 1, complete					

ORGANISM
Holo sapientis
Eukaryota; Metazoa; Chordata; Craniata; Vertebrata; Euteleostomi;
Mammalia; Platyrrhini; Hominoidea; Hominidae; Homo

AUTHORS	Williams, S.
TITLE	Direct Submission

COMMENTS
requests: clonerequest@sanger.ac.uk
On or before May 15 2001 this sequence version ren[er]ed

On 24 Dec 2001, the sequence version reported
gi:13160267.
gi:7283183.
During sequence assembly data is compared from overlapping clones.
Where differences are found these are annotated as variations
between the clones of the same library. Note that the

together with a note of the overlapping clone name. Note that the variation annotation may not be found in the sequence submission corresponding to the overlapping clone, as we submit sequences with only a small overlap as described above.

This sequence was finished as follows unless otherwise noted: all regions were either double-stranded or sequenced with an alternate chemistry or covered by high quality data (i.e., phred quality >= 30): an attempt was made to resolve all sequencing problems, such

assembly was confirmed by restriction digest. The following

abbreviations are used to associate primary accession numbers given in the feature table with their source databases: Em, EMBL; Sw, SWISSPROT; Tr, TREMBL; Wp, WORMPEP; Information on the WORMPEP database can be found at

http://www.sanger.ac.uk/Projects/C_elegans/wormmap This sequence was generated from part of bacterial clone contigs of human chromosome 1, constructed by the Sanger Centre Chromosome 1 Mapping Group. Further information can be found at

RP11-3J23 is from the library RPCI-11.1 constructed by the group of Pieter de Jong. For further details see <http://www.sanger.ac.uk/HGP/Chr1>

http://www.chori.org/bacpac/home.htm
VECTOR: pBACE3.6
IMPORTANT: This sequence is not the entire insert of clone RP11-3J23. It may be shorter because we sequence overlapping

sections only once, except for a 100 base overlap. The true left end of clone RP4-63318 is at 103414 in this sequence. The true right end of clone RP5-115A15 is at 100 in this sequence.

FEATURES	LOCATION/VALUES
SOURCE	1. 103513 /organism="Homo sapiens" /mol_type="genomic DNA"

```

/db_xref="caxon:9606"
/chromosome="1"
/clone="RP11-3723"
/clone_1fb="RP11-11.1"

```

```
repeat_region      870. .941
                    /note="36 copies 2 mer aa 65% conserved"
repeat_region      877. .921
                    /note="36 copies 2 mer aa 65% conserved"
```

```

repeat_region      1178.  .1469      /notes="5 copies 9 mer atcaatada 66% conserved"
repeat_region      1477.  .1752      /notes="146 copies 2 mer aa 55% conserved"

```

```

repeat_region /note="AluJo repeat: matches 3. .285 of consensus"
repeat_region /note="AluJo repeat: matches 3. .2365 of consensus"
repeat_region /note="AluJo repeat: matches 3. .300 of consensus"
repeat_region /note="AluJo repeat: matches 3. .3429 of consensus"

```



```

Fragment Name      Begin      End
LMFLCHR36_00      1      110000
LMFLCHR36_01      100001      210000
LMFLCHR36_02      200001      310000
LMFLCHR36_03      300001      410000
LMFLCHR36_04      400001      510000
LMFLCHR36_05      500001      610000
LMFLCHR36_06      600001      710000
LMFLCHR36_07      700001      810000
LMFLCHR36_08      800001      910000
LMFLCHR36_09      900001      1010000
LMFLCHR36_10      1000001      1110000
LMFLCHR36_11      1100001      1210000
LMFLCHR36_12      1200001      1310000
LMFLCHR36_13      1300001      1410000
LMFLCHR36_14      1400001      1510000
LMFLCHR36_15      1500001      1610000
LMFLCHR36_16      1600001      1710000
LMFLCHR36_17      1700001      1810000
LMFLCHR36_18      1800001      1910000
LMFLCHR36_19      1900001      2010000
LMFLCHR36_20      2000001      2110000
LMFLCHR36_21      2100001      2210000
LMFLCHR36_22      2200001      2310000
LMFLCHR36_23      2300001      2410000
LMFLCHR36_24      2400001      2510000
LMFLCHR36_25      2500001      2610000
LMFLCHR36_26      2600001      2710000
LMFLCHR36_27      2700001      2810000
LMFLCHR36_28      2800001      2910000
LMFLCHR36_29      2900001      3010000
LMFLCHR36_30      3000001      3110000
LMFLCHR36_31      3100001      3210000
LMFLCHR36_32      3200001      3310000
LMFLCHR36_33      3300001      3410000
LMFLCHR36_34      3400001      3510000
LMFLCHR36_35      3500001      359882
Continuation (12 of 36) of LMFLCHR36 from base 1100001 (AL499624 Letismania major chromo

```

```

Query Match      27.7%; Score 18; DB 2; Length 110000;
Best Local Similarity 100.0%; Pred. No. 19;
Matches 18; Conservative 0; Mismatches 0; Indels 0; Gaps 0;

```

```

QY      14 CTTCCGCTTTTCACGCG 31
          |||||
Db      66513 CTTCCGCTTTTCACGCG 66496

```

```

RESULT 10      AC079838      156481 bp      DNA      linear      HTG 01-APR-2001
LOCUS      AC079838
DEFINITION      Homo sapiens chromosome 2 clone RP11-257N14, WORKING DRAFT
ACCESSION      AC079838
VERSION      AC079838.3 GI:13493151
KEYWORDS      HTG; HTGS_PHASE1; HTGS_DRAFT; HTGS_FULLTOP.
SOURCE      Homo sapiens (human)
ORGANISM      Homo sapiens
Eukaryota; Metazoa; Chordata; Craniata; Vertebrata; Euteleostomi;
Mammalia; Eutheria; Primates; Catarrhini; Homnidae; Homo.
1 (bases 1 to 156481)
Waterston,R.H.
The sequence of Homo sapiens clone
Unpublished
2 (bases 1 to 156481)
Waterston,R.H.
Direct Submission
Submitted (12-SEP-2000) Genome Sequencing Center, Washington
University School of Medicine, 444 Forest Park Parkway, St. Louis,
MO 63108, USA
On Apr 1, 2001 this sequence version replaced gi:10799482.
COMMENT
----- Genome Center -----

```

```

Center: Washington University Genome Sequencing Center
Center code: WUGSC
Web site: http://genome.wustl.edu/gsc/index.shtml
----- Project Information -----
Center project name: H.NH0257N14
----- Summary Statistics -----
Sequencing vector: M13; 68%
Sequencing vector: plasmid; 32%
Chemistry: Dye-primer ET; 68% of reads
Chemistry: Dye-terminator Big Dye; 32% of reads
Assembly program: Phrap; version 0.99019
Consensus quality: 154814 bases at least Q40
Consensus quality: 155806 bases at least Q30
Insert size: 139000; agarose-fp
Insert size: 156381; sum-of-contigs
Quality coverage: 6.40 in Q20 bases; agarose-fp
Quality coverage: 6.28 in Q20 bases; sum-of-contigs
-----
* NOTE: This is a 'working draft' sequence. It currently
* consists of 2 contigs. The true order of the pieces
* is not known and their order in this sequence record is
* arbitrary. Gaps between the contigs are represented as
* runs of N, but the exact sizes of the gaps are unknown.
* This record will be updated with the finished sequence
* as soon as it is available and the accession number will
* be preserved.
*
* 1      6178: contig of 6178 bp in length
*
* 6179      6278: gap of unknown length
*
* 6279      156481: contig of 150203 bp in length.
Location/Qualifiers
1. 156481
/organism="Homo sapiens"
/mol_type="genomic DNA"
/db_xref="taxon:9606"
/chromosome="2"
/clone="RP11-257N14"
1. 6178
/note="assembly_name:Contig7
clone_end:17
vector_side:right"
6279. 156481
/note="assembly_name:Contig8
clone_end:SP6
vector_side:left"

```

```

misc_feature
/note="assembly_name:Contig7
clone_end:17
vector_side:right"
6279. 156481
/note="assembly_name:Contig8
clone_end:SP6
vector_side:left"
misc_feature

```

```

Query Match      27.7%; Score 18; DB 2; Length 156481;
Best Local Similarity 100.0%; Pred. No. 18;
Matches 18; Conservative 0; Mismatches 0; Indels 0; Gaps 0;

```

```

QY      43 CCTGCTGATGACACCC 60
          |||||
Db      11249 CCTGCTGATGACACCC 11266

```

```

RESULT 11      AC007098      157650 bp      DNA      linear      PRI 09-MAY-2001
LOCUS      AC007098
DEFINITION      Homo sapiens BAC clone RP11-435f16 from 2, complete sequence.
ACCESSION      AC007098
VERSION      AC007098.4 GI:11038585
KEYWORDS      HTG.
SOURCE      Homo sapiens (human)
ORGANISM      Homo sapiens
Eukaryota; Metazoa; Chordata; Craniata; Vertebrata; Euteleostomi;
Mammalia; Eutheria; Primates; Catarrhini; Homnidae; Homo.
1 (bases 1 to 157650)
Sulston,J.B. and Waterston,R.
Toward a complete human genome sequence
Genome Res. 8 (11), 1097-1108 (1998)
99063792
MEDLINE
PUBMED
9847074

```

REFERENCE 2 (bases 1 to 157650)
 AUTHORS Kallunki, J., Johnson, D. and Harris, A.
 TITLE The sequence of Homo sapiens BAC clone RP11-443F16
 JOURNAL Unpublished
 REFERENCE 3 (bases 1 to 157650)
 AUTHORS Waterston, R.H.
 TITLE Direct Submission
 JOURNAL Submitted (16-MAR-1999) Genome Sequencing Center, Washington University School of Medicine, 4444 Forest Park Parkway, St. Louis, MO 63108, USA

REFERENCE 4 (bases 1 to 157650)
 AUTHORS Waterston, R.H.
 TITLE Direct Submission
 JOURNAL Submitted (30-OCT-2000) Genome Sequencing Center, Washington University School of Medicine, 4444 Forest Park Parkway, St. Louis, MO 63108, USA

REFERENCE 5 (bases 1 to 157650)
 AUTHORS Waterston, R.
 TITLE Direct Submission
 JOURNAL Submitted (09-MAY-2001) Department of Genetics, Washington University, 4444 Forest Park Avenue, St. Louis, Missouri 63108, USA
 On Oct 30, 2000 this sequence version replaced gi:8570191.

COMMENT
 Center: Washington University Genome Sequencing Center
 Center code: WUGSC
 Web site: <http://genome.wustl.edu/gsc>
 Contact: sapient@wustl.wustl.edu
 Summary Statistics
 Center project name: H_NH0443F16

NOTICE: This sequence may not represent the entire insert of this clone. It may be shorter because we only sequence overlapping clone sections once, or longer because we provide a small overlap between neighboring data submissions.

This sequence was finished as follows unless otherwise noted:
 all regions were double stranded, sequenced with an alternate chemistry, or covered by high quality data (i.e., phred quality >= 30); an attempt was made to resolve all sequencing problems, such as compressions and repeats; all regions were covered by sequence from more than one subclone; and the assembly was confirmed by restriction digest.

MAPPING INFORMATION:
 Mapping information for this clone was provided by Dr. John D. McPherson, Department of Genetics, Washington University, St. Louis MO. For additional information about the map position of this sequence, see <http://genome.wustl.edu/gsc>

SOURCE INFORMATION:
 The RFL1-11 human BAC library was made from the blood of one male donor, as described by Osogawa, K., Moon, P.Y., Zhao, B., Frenken, E., Tateo, M., Catanesi, J.D. and de Jong, P.J. (1998) An improved approach for construction of bacterial artificial chromosome libraries. Genomics 51:1-8. The clone may be obtained either from Research Genetics, Inc. (<http://www.rgsen.com>) or Pieter de Jong and coworkers at the Roswell Park Cancer Institute (<http://bacpac.med.buffalo.edu>)
 VECTOR: pBAC3.6

NEIGHBORING SEQUENCE INFORMATION:
 The clone sequenced to the right is RP11-511I11, 200 bp overlap. Actual start of this clone is at base position 1 of RP11-443F16; actual end is at base position 157456 of RP11-443F16.

FEATURES
 source
 1. 157650
 /organism="Homo sapiens"
 /mol_type="genomic DNA"
 /db_xref="taxon:9606"
 /chromosome="2"
 /map="2"
 /clone="RP11-443F16"
 /clone_1bp="RP11-11"

misc_feature 75..449
 /note="similar to EST H66784 (NID:g1025524) YR84h10.r1"
 repeat_region 181..425
 /rpt_family="L1"
 repeat_region 422..1148
 /rpt_family="L1"
 repeat_region 1149..1476
 /rpt_family="Alu"
 repeat_region 1477..1870
 /rpt_family="L1"
 repeat_region 1871..2176
 /rpt_family="Alu"
 repeat_region 2177..2455
 /rpt_family="L1"
 repeat_region 2228..2698
 /rpt_family="L1"
 repeat_region 2800..3357
 /rpt_family="MALR"
 repeat_region 3497..3582
 /rpt_family="L2"
 repeat_region 3599..3949
 /rpt_family="L2"
 repeat_region 4875..4907
 /rpt_family="L2"
 repeat_region 5004..5103
 /rpt_family="L2"
 repeat_region 5348..5454
 /rpt_family="L2"
 repeat_region 6944..7100
 /rpt_family="MER2_type"
 repeat_region 7099..7878
 /rpt_family="MER2_type"
 repeat_region 7912..8544
 /rpt_family="MER2_type"
 repeat_region 8545..10505
 /rpt_family="ERV1"
 repeat_region 10510..10614
 /rpt_family="MER2_type"
 repeat_region 10619..10995
 /rpt_family="MALR"
 repeat_region 11078..11306
 /rpt_family="L2"
 misc_feature 12958..13086
 /note="similar to EST AA456682 (NID:g2179258) aal3e01.r1"
 misc_feature 12958..13086
 /note="similar to EST A1150782 (NID:g3679251) qc06g03.x1"
 misc_feature 12958..13086
 /note="similar to EST A1675083 (NID:g4875563) wc23c03.x1"
 misc_feature 12965..13086
 /note="similar to EST AA721648 (NID:g2736631) ny87f02.s1"
 misc_feature 13045..13086
 /note="similar to EST AM102800 (NID:g6073413) xd38c10.x1"
 repeat_region 14003..14303
 /rpt_family="Alu"
 repeat_region 14920..15821
 /rpt_family="Alu"
 repeat_region 15257..15864
 /rpt_family="Alu"
 repeat_region 16132..16425
 /rpt_family="Alu"
 repeat_region 16796..17077
 /rpt_family="Alu"
 repeat_region 17109..17251
 /rpt_family="Alu"
 repeat_region 17636..17941
 /rpt_family="Alu"
 repeat_region 17948..18103
 /rpt_family="Alu"
 misc_feature 18324..18766
 /note="similar to EST Z21249 (NID:g279922)"
 misc_feature 18526..18766
 /note="similar to EST AM102800 (NID:g6073413) xd38c10.x1"
 misc_feature 18526..18699

```

misc_feature      /note="similar to EST A1675083 (NID:g4875563) wc23c03.x1"
18526..18691
misc_feature      /note="similar to EST AA721648 (NID:g2736631) ny87f02.s1"
18526..18691
misc_feature      /note="similar to EST A1150782 (NID:g3679251) qc06g03.x1"
18526..18691
misc_feature      /note="similar to EST AA456682 (NID:g2179258) aa13e01.r1"
18583..18780
misc_feature      /note="similar to EST A1806986 (NID:g5393552) wf24g10.x1"
19222..19264
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Matches 18; Conservative 0; Mismatches 0; Indels 0; Gaps 0;

QY 43 CCTGCTGATGACACCC 60
DB 28647 CCTGCTGATGACACCC 28664

RESULT 12
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LOCUS Homo sapiens clone NH0443B15a, *** SEQUENCING IN PROGRESS ***, 6
DEFINITION
AC007629
ACCESSION
AC007629
VERSION
AC007629.9 GI:10716584
KEYWORDS
HTG; HTGS PHASE1.
SOURCE
Homo sapiens (human)
ORGANISM
Homo sapiens
Eukaryota; Metazoa; Chordata; Craniata; Vertebrata; Euteleostomi;
Mammalia; Eutheria; Primates; Catarrhini; Homiidae; Homo.
REFERENCE
1 (bases 1 to 157711)
AUTHORS
McCombie, W.R.
TITLE
Human Genomic Sequence
JOURNAL
Unpublished
REFERENCE
2 (bases 1 to 157711)
AUTHORS
McCombie, W.R.
TITLE
Direct Submission
JOURNAL
Submitted (21-MAY-1999) Lite Annenberg Hazen Genome Sequencing
Center, Cold Spring Harbor Laboratory, 1 Bungtown Road, Cold Spring
Harbor, NY 11724, USA
On Oct 7, 2000 this sequence version replaced gi:9972279.
* NOTE: This is a 'working draft' sequence. It currently

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* consists of 6 contigs. The true order of the pieces
* is not known and their order in this sequence record is
* arbitrary. Gaps between the contigs are represented as
* runs of N, but the exact sizes of the gaps are unknown.
* This record will be updated with the finished sequence
* as soon as it is available and the accession number will
* be preserved.
* 1 142198: contig of 142198 bp in length
* 142199 142298: gap of unknown length
* 142299 157945: contig of 15647 bp in length
* 157945 158045: gap of unknown length
* 158045 160827: contig of 2782 bp in length
* 160828 160928: gap of unknown length
* 160928 163434: contig of 2507 bp in length
* 163435 163534: gap of unknown length
* 163535 165854: contig of 2320 bp in length
* 165855 165954: gap of unknown length
* 165955 167711: contig of 1757 bp in length.

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ORIGIN
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Best Local Similarity 100.0%; Pred. No. 18;
Matches 18; Conservative 0; Mismatches 0; Indels 0; Gaps 0;

QY 43 CCTGCTGATGACACCC 60
DB 128804 CCTGCTGATGACACCC 128787

RESULT 13
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LOCUS Haemophilus ducreyi strain 35000HP section 6 of 6 of the complete
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DEFINITION
AE017156 AE017143
ACCESSION
AE017156
VERSION
AE017156.1 GI:33149035
KEYWORDS
Haemophilus ducreyi 35000HP
SOURCE
Haemophilus ducreyi 35000HP
Bacteria; Proteobacteria; Gammaproteobacteria; Pasteurellales;
Pasteurellaceae; Haemophilus.
REFERENCE
1 (bases 1 to 191854)
AUTHORS
Munson, R.S., Jr., Ray, W.C., Mahairas, G., Sabo, P., Mungur, R.,
Johnson, L., Nguyen, D., Wang, J., Forst, C. and Hood, L.
TITLE
The Complete Genome Sequence of Haemophilus ducreyi
JOURNAL
Unpublished
2 (bases 1 to 191854)
AUTHORS
Munson, R.S., Jr., Ray, W.C., Mahairas, G., Sabo, P., Mungur, R.,
Johnson, L., Nguyen, D., Wang, J., Forst, C. and Hood, L.
TITLE
Direct Submission
JOURNAL
Submitted (04-JUN-2003) Pediatrics, Columbus Children's Research
Institute and The Ohio State University, 700 Children's Drive,
Columbus, OH 43205 USA

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Best Local Similarity 100.0% Pred. No. 18;
Matches 18; Conservative 0; Mismatches 0; Indels 0; Gaps 0;
QY 40 AAACCTGCTGATGACA 57
Db 119863 AAACCTGCTGATGACA 119860

RESULT 14
LOCUS AE016888
DEFINITION Eremothecium gossypii chromosome III, section 3 of 3 of the
complete sequence.
ACCESSION AE016888
VERSION AE016888.1
KEYWORDS GI:44981618
SOURCE Eremothecium gossypii (Ashbya gossypii)
ORGANISM Eremothecium gossypii
Eukaryota; Fungi; Ascomycota; Saccharomycotina; Saccharomycetes;
Saccharomycetales; Saccharomycetaceae; Eremothecium.
REFERENCE 1 (bases 1 to 306790)
Dietrich, F.S., Voegelé, S., Brachat, S., Lerch, A., Gates, K.,
Steiner, S., Mohr, C., Pohlmann, R., Luedi, P., Choi, S., Wing, R.A.,
Flavler, A., Gaffney, T.D. and Philippsen, P.
The Ashbya gossypii genome as a tool for mapping the ancient
Saccharomyces cerevisiae genome
Science 304 (5668), 304-307 (2004)
JOURNAL 15001715
PUBMED 2 (bases 1 to 306790)
Brachat, S., Voegelé, S.E., Dietrich, F.S., Lerch, A., Gaffney, T. and
Philippsen, P.
Direct Submision
Submitted (20-DEC-2002) Applied Microbiology, Biozentrum,
University of Basel, Klingelbergstrasse 50-70, Basel CH-4056,
Switzerland
COMMENT This is low coverage sequence generated to identify the complete
set of genes and the gene order on this chromosome. Regions of low
quality are identified. Before doing extensive work on any gene
identified here the sequence should be confirmed.
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YPL154C (PEP4); tandem gene duplication in Ashbya
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LRAELALQHFENVNLDMVALPGOTRERADLETICNSRVASHVHLIENPTFF
YRPPQLPDESDADMOVNIHILIAERGVYRYSAPVQPKPLNNMYWXYGDYLG
ICAGASKTSIFRDRIRIOMRKHHPROYEQAAMACIENLATECEVQSDRIPEFM
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6979. .8820
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Matches 18; Conservative 0; Mismatches 0; Indels 0; Gaps 0;

OY 37 TCGAACCCTGCTGATGG 54
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Search completed: December 22, 2004, 23:36:25
Job time : 929.412 secs

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OM nucleic - nucleic search, using sw model

Run on: December 22, 2004, 19:14:28 ; Search time 229.412 Seconds

Title: US-09-898-616A-1

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Scoring table: OLIGO_NUC

Searched: 4134886 seqs, 2624710521 residues

Word size :

Total number of hits satisfying chosen parameters: 8269772

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Maximum DB seq length: 2000000000
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Post-processing: Listing first 45 summaries

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Pred. No. is the number of results predicted by chance to have a score greater than or equal to the score of the result being printed and is derived by analysis of the total score distribution.

SUMMARIES

Result No.	Score	Query Match	Length	DB	ID	Description
1	65	100.0	65	9	ABZ58370	AbZ58370 Human utt
2	65	100.0	65	12	ADL27626	ADL27626 Recombina
3	65	35.4	60	9	ABZ58376	AbZ58376 Human utt
4	23	35.4	60	9	ABZ58377	AbZ58377 Human utt
5	23	35.4	60	12	ADL27633	ADL27633 Recombina
6	23	35.4	60	12	ADL27632	ADL27632 Recombina
7	17	26.2	36	6	ABK10937	ABK10937 primer 1R
8	17	26.2	36	6	ABK74610	ABK74610 Interleuk
9	17	26.2	990	5	AAH66339	AAH66339 C glutami
10	17	26.2	1113	4	AAFP1853	AAFP1853 Corynebac
11	17	26.2	1113	4	AA596115	AA596115 C. glutam
12	17	26.2	2055	5	AA574960	AA574960 DNA encod
13	17	26.2	2877	8	ACAA3968	ACAA3968 Prokaryot
14	17	26.2	2892	4	AA544085	AA544085 Pseudomon
15	17	26.2	2892	8	ACAA42199	ACAA42199 Prokaryot
16	17	26.2	2932	8	ACA45765	ACA45765 Prokaryot
17	17	26.2	2987	11	ABPD0487	ABPD0487 Pseudomon
18	17	26.2	3009	11	ABPD04755	ABPD04755 Pseudomon
19	17	26.2	7642	10	ADD44755	ADD44755 Human gen
20	17	26.2	349960	5	AAH68528	AAH68528 C. glutami
21	16	24.6	139	12	ACH90361	ACH90361 Human gen

C	22	16	24.6	230	8	ABZ19128	Group	III
C	23	16	24.6	277	8	ABZ18853	Group	III
C	24	16	24.6	401	4	AAK95547	Human	neu
C	25	16	24.6	401	4	AAK97040	Human	neu
C	26	16	24.6	401	6	ABT00317	Human	neu
C	27	16	24.6	401	6	ABT01810	Human	neu
C	28	16	24.6	426	8	ACA43852	Prokaryot	
C	29	16	24.6	465	8	ABZ20021	Group	III
C	30	16	24.6	490	8	ABZ19219	Group	III
C	31	16	24.6	513	12	ACH76661	Human	ger
C	32	16	24.6	518	8	ABZ19817	Group	III
C	33	16	24.6	543	6	ABK73249	Bacilli	
C	34	16	24.6	629	8	ABZ19560	Group	III
C	35	16	24.6	646	8	ABZ19299	Group	III
C	36	16	24.6	894	11	ACH95360	Klebsiell	
C	37	16	24.6	1044	11	ABZ73819	Secreted	
C	38	16	24.6	1044	10	ADC20635	Human	sec
C	39	16	24.6	1044	10	ABT16890	Human	sec
C	40	16	24.6	1044	10	ABZ67403	Human	sec
C	41	16	24.6	1052	3	AAK59250	Human	sec
C	42	16	24.6	1052	8	ABZ73338	Secreted	
C	43	16	24.6	1052	10	ADC20039	Human	sec
C	44	16	24.6	1052	10	ABT16781	Human	sec
C	45	16	24.6	1052	10	ABZ65948	Human	sec

ALIGNMENTS

CC	Human uteroglobin synthetic gene oligonucleotide 1.
CC	Human; uteroglobin; respiratory distress; antiinflammatory; antifibrotic;
KM	antiinflammatory; antiasthmatic; nephrotropic; antipneumatic;
KM	antiarthritic; ss.
OS	Homo sapiens.
OS	Synthetic.
FN	WO2003003979-A2.
PD	16-JAN-2003.
PF	02-JUL-2002; 2002WO-US020836.
PR	02-JUL-2001; 2001US-00898616.
PA	(CLAR-) CLARAGEN INC.
PI	Pilon AL, Welch RE;
DR	WPI; 2003-221527/21.
XX	Bacterial expression system for producing recombinant human uteroglobin
XX	for treating inflammatory and fibrotic conditions, comprises a synthetic
XX	gene which codes for human uteroglobin.
XX	Claim 1; Page 33; 127bp; English.
XX	The present sequence is that of oligonucleotide 1, which was used in the
XX	construction of a synthetic gene for the production of human uteroglobin
XX	(hUG) in bacteria. Oligonucleotides 1-4 (see AB258370-73) were used to
XX	assemble the coding strand and oligonucleotides 5-8 (see AB258374-77) the
XX	complementary strand. The gene was assembled by annealing and ligation of
XX	the oligonucleotides. Because mature native hUG has glutamic acid at its
XX	N-terminus, an initiator methionine was added to the N-terminus, and

CC codon usage was optimised for expression in bacteria. In an example from
CC the invention, the synthetic gene was cloned into plasmid pCG12 (see
CC AB583378) and recombinant rhug (see ABP72259) was produced in *Escherichia*
CC coli strain CG12. The invention relates generally to the production of
CC recombinant rhug by bacterial expression, protein purification and scaled-
CC up production according to current good manufacturing practices. The
CC recombinant rhug is useful for the treatment of inflammatory and fibrotic
CC conditions, such as neonatal respiratory distress syndrome and
CC bronchopulmonary dysplasia. It may also be used to treat conditions
CC associated with elevated phospholipase A2 levels such as pancreatitis,
CC acute renal failure, rheumatoid arthritis and asthma
CC
SQ Sequence 65 BP; 13 A; 22 C; 14 G; 16 T; 0 U; 0 Other;
Query Match 100.0%; Score 65; DB 9; Length 65;
Best Local Similarity 100.0%; Pred. No. 2e-25;
Matches 65; Conservative 0; Mismatches 0; Indels 0; Gaps 0;
QY 1 GATCCATGGAATCTGCCGCTTTCCAGCGCTTATCGAAACCCCTGATGACACCC 60
Db 1 GATCCATGGAATCTGCCGCTTTCCAGCGCTTATCGAAACCCCTGATGACACCC 60
QY 61 CGTCC 65
Db 61 CGTCC 65
RESULT 2
ADL27626
ID ADL27626 standard; DNA; 65 BP.
XX ADL27626;
AC 20-MAY-2004 (first entry)
DT
XX
DE Recombinant human uteroglobin, rhug, coding oligonucleotide #1.
XX
XX Human; ss; recombinant human uteroglobin; rhug;
XX Bacterial expression system; rhug master cell bank;
XX rhug research seed bank; anti-inflammatory; secretory phospholipase A 2;
XX fibronectin; respiratory distress; inflammation; fibrotic disease.
XX
XX Homo sapiens.
OS Synthetic.
OS
XX US2003207795-A1.
XX
XX 06-NOV-2003.
PD
XX
XX 02-JUL-2002; 2002US-00187498.
PF
XX 28-MAY-1997; 97US-00864357.
PR
XX 02-JUL-2001; 2001US-00898616.
XX
XX (PILIO/) PILON A L.
PA (WELC/) WELCH R W.
XX
XX Pilon AL, Welch RW;
PI
XX
XX WPI, 2004-051527/05.
DR
XX
XX Bacterial expression system for production of recombinant human
PT uteroglobin comprising synthetic gene or human cDNA sequence which codes
PT for human uteroglobin.
XX
XX Claim 1; SEQ ID NO 1; 64bp; English.
PS
XX The invention relates to a bacterial expression system for the production
CC of recombinant human uteroglobin (rhug), comprising a synthetic gene or
CC human cDNA sequence which codes for human rhug, constructed from the
CC oligonucleotides appearing as ADL27626-ADL27629, and which further
CC comprises Met-Ala-Ala at the N-terminus of the sequence. Also included
CC are producing an rhug master cell bank (comprising inoculating a suitable

CC incubating broth with an aliquot portion of a rhug research seed bank to
CC form a bacterial culture, incubating the bacterial culture, adding a
CC cryoprotective to the bacterial culture to form a cryopreserved
CC solution, transferring a portion of the cryopreserved solution to a
CC cryovial and storing the cryovial at a temperature below -60 degrees C).
CC expressing rhug (comprising providing a production seed cell bank culture
CC comprising an expression vector capable of expressing rhug, inoculating a
CC broth medium with the production seed cell bank culture to form an
CC inoculum, incubating the bacterial culture formed in step (b),
CC inoculating a large scale fermenter with the inoculum formed from the
CC step (c) to form a fermentation culture, incubating the fermentation
CC culture within the large scale fermenter, adding an induction agent to
CC the fermentation culture to induce the expression of rhug and harvesting
CC of rhug in a sample, measuring in vitro anti-inflammatory effect arising
CC from inhibition or blocking of secretory phospholipase A 2 enzymes by
CC rhug, measuring in vitro binding of rhug to fibronectin, determining the
CC purity of rhug, and a pharmaceutical composition comprising a purified
CC rhug and a carrier or diluent. The bacterial expression system is useful
CC for producing a rhug research seed bank or a pharmaceutical grade rhug
CC drug substance. rhug is safe to administer to a patient in respiratory
CC distress. The rhug is useful for treating inflammation and fibrotic
CC diseases. The present sequence is a coding strand oligonucleotide used to
CC construct the synthetic rhug gene.
XX
SQ Sequence 65 BP; 13 A; 22 C; 14 G; 16 T; 0 U; 0 Other;
Query Match 100.0%; Score 65; DB 12; Length 65;
Best Local Similarity 100.0%; Pred. No. 2e-25;
Matches 65; Conservative 0; Mismatches 0; Indels 0; Gaps 0;
QY 1 GATCCATGGAATCTGCCGCTTTCCAGCGCTTATCGAAACCCCTGATGACACCC 60
Db 1 GATCCATGGAATCTGCCGCTTTCCAGCGCTTATCGAAACCCCTGATGACACCC 60
QY 61 CGTCC 65
Db 61 CGTCC 65
RESULT 3
ABZ58376/c
ID ABZ58376 standard; DNA; 60 BP.
XX ABZ58376;
AC 28-APR-2003 (first entry)
DT
XX
XX Human uteroglobin synthetic gene oligonucleotide 7.
DE
XX
XX Human; uteroglobin; respiratory distress; anti-inflammatory; antifibrotic;
XX anti-inflammatory; antiaesthetic; nephroretropic; antirheumatic;
XX antiarthritic; ss.
XX
XX Homo sapiens.
OS Synthetic.
OS
XX WO2003003979-A2.
XX
XX 16-JAN-2003.
PD
XX
XX 02-JUL-2002; 2002WO-US020836.
PF
XX 02-JUL-2001; 2001US-00898616.
PR
XX
XX (CLAR-) CLARAGEN INC.
PA
XX
XX Pilon AL, Welch RE;
PI
XX
XX WPI, 2003-221527/21.
DR
XX Bacterial expression system for producing recombinant human uteroglobin
PT for treating inflammatory and fibrotic conditions, comprises a synthetic

PT Gene which codes for human uteroglobin.
XX
PS Example 1; Page 33; 127pp; English.
XX
CC The present sequence is that of oligonucleotide 7, which was used in the
CC construction of a synthetic gene for the production of human uteroglobin
CC (hUG) in bacteria. Oligonucleotides 1-4 (see AB258370-73) were used to
CC assemble the coding strand and oligonucleotides 5-8 (see AB258374-77) the
CC complementary strand. The gene was assembled by annealing and ligation of
CC the oligonucleotides. Because mature native hUG has glutamic acid at its
CC N-terminus, an initiator methionine was added to the N-terminus, and
CC codon usage was optimised for expression in bacteria. In an example from
CC the invention, the synthetic gene was cloned into plasmid pCG12 (see
CC AB258378) and recombinant hUG (see ABP72259) was produced in *Escherichia*
CC coli strain CG12. The invention relates generally to the production of
CC recombinant hUG by bacterial expression, protein purification and scaled-
CC up production according to current good manufacturing practices. The
CC recombinant hUG is useful for the treatment of inflammatory and fibrotic
CC conditions, such as neonatal respiratory distress syndrome and
CC bronchopulmonary dysplasia. It may also be used to treat conditions
CC associated with elevated phospholipase A2 levels such as pancreatitis,
CC acute renal failure, rheumatoid arthritis and asthma
XX
SQ Sequence 60 BP; 12 A; 14 C; 22 G; 12 T; 0 U; 0 Other;
Query Match 35.4%; Score 23; DB 9; Length 60;
Best Local Similarity 100.0%; Pred. No. 0.014;
Matches 23; Conservative 0; Mismatches 0; Indels 0; Gaps 0;
XX
QY 43 CCTGCTGATGACACCCCGTCC 65
DB 60 CCTGCTGATGACACCCCGTCC 38
RESULT 4
AB258377/c
ID AB258377 standard; DNA; 60 BP.
XX
AC AB258377;
XX
DT 28-APR-2003 (first entry)
XX
DE Human uteroglobin synthetic gene oligonucleotide 8.
XX
KW Human; uteroglobin; respiratory distress; antiinflammatory; antifibrotic;
KW antiinflammatory; antiasthmatic; nephrotropic; antirheumatic;
XX
XX Homo sapiens.
OS
XX Synthetic.
XX
XX WO2003003979-A2.
XX
XX 16-JAN-2003.
XX
XX 02-JUL-2002; 2002WO-US020836.
XX
XX 02-JUL-2001; 2001US-00898616.
XX
XX (CLAR-) CLARAGEN INC.
XX
XX Pilon AL, Welch RE;
XX
XX WPI; 2003-221527/21.
XX
XX Bacterial expression system for producing recombinant human uteroglobin
PT for treating inflammatory and fibrotic conditions, comprises a synthetic
PT gene which codes for human uteroglobin.
XX
XX Example 1; Page 33; 127pp; English.
XX
XX The present sequence is that of oligonucleotide 8, which was used in the
CC construction of a synthetic gene for the production of human uteroglobin

CC (hUG) in bacteria. Oligonucleotides 1-4 (see AB258370-73) were used to
CC assemble the coding strand and oligonucleotides 5-8 (see AB258374-77) the
CC complementary strand. The gene was assembled by annealing and ligation of
CC the oligonucleotides. Because mature native hUG has glutamic acid at its
CC N-terminus, an initiator methionine was added to the N-terminus, and
CC codon usage was optimised for expression in bacteria. In an example from
CC the invention, the synthetic gene was cloned into plasmid pCG12 (see
CC AB258378) and recombinant hUG (see ABP72259) was produced in *Escherichia*
CC coli strain CG12. The invention relates generally to the production of
CC recombinant hUG by bacterial expression, protein purification and scaled-
CC up production according to current good manufacturing practices. The
CC recombinant hUG is useful for the treatment of inflammatory and fibrotic
CC conditions, such as neonatal respiratory distress syndrome and
CC bronchopulmonary dysplasia. It may also be used to treat conditions
CC associated with elevated phospholipase A2 levels such as pancreatitis,
CC acute renal failure, rheumatoid arthritis and asthma
XX
SQ Sequence 60 BP; 12 A; 14 C; 22 G; 12 T; 0 U; 0 Other;
Query Match 35.4%; Score 23; DB 9; Length 60;
Best Local Similarity 100.0%; Pred. No. 0.014;
Matches 23; Conservative 0; Mismatches 0; Indels 0; Gaps 0;
XX
QY 43 CCTGCTGATGACACCCCGTCC 65
DB 60 CCTGCTGATGACACCCCGTCC 38
RESULT 5
ADL27633/c
ID ADL27633 standard; DNA; 60 BP.
XX
AC ADL27633;
XX
DT 20-MAY-2004 (first entry)
XX
DE Recombinant human uteroglobin, rhUG, non-coding oligonucleotide #4.
XX
XX Human; sg; recombinant human uteroglobin; rhUG;
KW bacterial expression system; rhUG master cell bank;
KW rhUG research seed bank; anti-inflammatory; secretory phospholipase A 2;
KW fibronectin; respiratory distress; inflammation; fibrotic disease.
XX
XX Homo sapiens.
OS
XX Synthetic.
XX
XX US2003207795-A1.
XX
XX 06-NOV-2003.
XX
XX 02-JUL-2002; 2002US-00187498.
XX
XX 28-MAY-1997; 97US-00864357.
XX
XX 02-JUL-2001; 2001US-00898616.
XX
XX (PILO/) PILON A L.
XX
XX (WELC/) WELCH R W.
XX
XX Pilon AL, Welch RW;
XX
XX WPI; 2004-051527/05.
XX
XX Bacterial expression system for production of recombinant human
PT uteroglobin comprising synthetic gene or human cDNA sequence which codes
PT for human uteroglobin.
XX
XX Example 1; SEQ ID NO 8; 64pp; English.
XX
XX The invention relates to a bacterial expression system for the production
CC of recombinant human uteroglobin (rhUG), comprising a synthetic gene or
CC human cDNA sequence which codes for human UG, constructed from the
CC oligonucleotides appearing as ADL27626-ADL27629, and which further
CC comprises Met-Ala-Ala at the N-terminus of the sequence. Also included

are producing an rhUG master cell bank (comprising inoculating a suitable incubating broth with an aliquot portion of a rhUG research seed bank to form a bacterial culture, incubating the bacterial culture, adding a cryopreservative to the bacterial culture to form a cryopreserved solution, transferring the cryopreserved solution to a cryovial and storing the cryovial at a temperature below -60 degrees C), expressing rhUG (comprising providing a production seed cell bank culture comprising an expression vector capable of expressing rhUG, inoculating a broth medium with the production seed cell bank culture to form an inoculum, incubating the bacterial culture formed in step (b), inoculating a large scale fermenter with the inoculum formed from the step (c) to form a fermentation culture, adding an induction agent to the fermentation culture to induce the expression of rhUG and harvesting the above fermentation culture), purifying rhUG, determining the potency of rhUG in a sample, measuring in vitro anti-inflammatory effect arising from inhibition or blocking of secretory phospholipase A 2 enzymes by rhUG, measuring in vitro binding of rhUG to fibronectin, determining the purity of rhUG, and a pharmaceutical composition comprising a purified rhUG and a carrier or diluent. The bacterial expression system is useful for producing a rhUG research seed bank or a pharmaceutical grade rhUG drug substance. rhUG is safe to administer to a patient in respiratory distress. The rhUG is useful for treating inflammation and fibrotic diseases. The present sequence is a non-coding strand oligonucleotide used to construct the synthetic rhUG gene.

Sequence 60 BP; 12 A; 14 C; 22 G; 12 T; 0 U; 0 Other;

Query Match 35.4%; Score 23; DB 12; Length 60;
Best Local Similarity 100.0%; Pred. No. 0.014;
Matches 23; Conservative 0; Mismatches 0; Indels 0; Gaps 0;

43 CCTGCTGATGACACCCCGTCC 65
60 CCTGCTGATGACACCCCGTCC 38

Db

RESULT 6
ADL27632/C
ID ADL27632 standard; DNA; 60 BP.

AC ADL27632;
XX
DT 20-MAY-2004 (first entry)
XX
DE Recombinant human uteroglobin, rhUG, non-coding oligonucleotide #3.
XX
KW Human; ss; recombinant human uteroglobin; rhUG;
KW bacterial expression system; rhUG master cell bank;
KW rhUG research seed bank; anti-inflammatory; secretory phospholipase A 2;
KW fibronectin; respiratory distress; inflammation; fibrotic disease.
XX
OS Homo sapiens.
OS Synthetic.
XX
PN US2003207795-A1.
XX
PD 06-NOV-2003.
XX
PF 02-JUL-2002; 2002US-00187498.
XX
PR 28-MAY-1997; 97US-00864357.
PR 02-JUL-2001; 2001US-00898616.
XX
XX (PILO/) PILON A L.
PA (WELC/) WELCH R W.
XX
PI Pilon AL, Welch RW;
XX
DR WPI, 2004-051527/05.
XX
PT Bacterial expression system for production of recombinant human uteroglobin comprising synthetic gene or human CDNA sequence which codes

for human uteroglobin.

XX
XX Example 1; SEQ ID NO 7; 64bp; English.

XX
XX The invention relates to a bacterial expression system for the production of recombinant human uteroglobin (rhUG), comprising a synthetic gene or human CDNA sequence which codes for human UG, constructed from the oligonucleotides appearing as ADL27626-ADL27629, and which further comprises Met-Ala-Ala at the N-terminus of the sequence. Also included are producing an rhUG master cell bank (comprising inoculating a suitable incubating broth with an aliquot portion of a rhUG research seed bank to form a bacterial culture, incubating the bacterial culture, adding a cryopreservative to the bacterial culture to form a cryopreserved solution, transferring the cryovial at a temperature below -60 degrees C), expressing rhUG (comprising providing a production seed cell bank culture comprising an expression vector capable of expressing rhUG, inoculating a broth medium with the production seed cell bank culture to form an inoculum, incubating the bacterial culture formed in step (b), inoculating a large scale fermenter with the inoculum formed from the step (c) to form a fermentation culture, adding an induction agent to the fermentation culture to induce the expression of rhUG and harvesting the above fermentation culture), purifying rhUG, determining the potency of rhUG in a sample, measuring in vitro anti-inflammatory effect arising from inhibition or blocking of secretory phospholipase A 2 enzymes by rhUG, measuring in vitro binding of rhUG to fibronectin, determining the purity of rhUG, and a pharmaceutical composition comprising a purified rhUG and a carrier or diluent. The bacterial expression system is useful for producing a rhUG research seed bank or a pharmaceutical grade rhUG drug substance. rhUG is safe to administer to a patient in respiratory distress. The rhUG is useful for treating inflammation and fibrotic diseases. The present sequence is a non-coding strand oligonucleotide used to construct the synthetic rhUG gene.

Sequence 60 BP; 12 A; 14 C; 22 G; 12 T; 0 U; 0 Other;

Query Match 35.4%; Score 23; DB 12; Length 60;
Best Local Similarity 100.0%; Pred. No. 0.014;
Matches 23; Conservative 0; Mismatches 0; Indels 0; Gaps 0;

43 CCTGCTGATGACACCCCGTCC 65
60 CCTGCTGATGACACCCCGTCC 38

Db

RESULT 7
ABK10937
ID ABK10937 standard; DNA; 36 BP.

AC ABK10937;
XX
DT 20-MAY-2002 (first entry)
XX
DE Primer IRAP5 relating to modified retroviral vector invention.
XX
KW Human; retrovirus vector; elongation factor 1-alpha; gene therapy;
KW BP 1alpha; Murine Leukemia Virus; MLV; multi-cloning site; MCS; primer;
KW IRAP5; ss.
XX
OS Unidentified.
OS
PN KR2001069245-A.
XX
PD 25-JUL-2001.
XX
PF 08-SEP-2000; 2000KR-00053613.
XX
PR 08-SEP-2000; 2000KR-00053613.
XX
XX (VIR0-) VIR0MED LTD.
PA
XX
PI Kim SY, Lee JT, Yoo SS;

XX WPI; 2002-065240/09.
XX Retroviral vectors useful in gene therapy, containing no viral coding
XX sequences but which includes a human elongation factor 1 alpha non-coding
XX fragment.
XX
XX Disclosure; Page 31; 32pp; Korean.
XX
XX The present invention relates to a retrovirus vector containing modified
XX non-coding sequences derived from the human elongation factor 1-alpha
XX gene (EF 1alpha, bases +773 to +1006), which can be effectively used in
XX gene therapy due to its high stability and expression. The Murine
XX Leukaemia Virus (MLV)-derived retrovirus vector is modified by complete
XX removal of the gag, env and pol genes to improve gene expression and
XX virus production. It is also modified to contain the non-coding
XX portion of the EF 1alpha gene upstream of a multi-cloning site (MCS). The
XX present sequence represents a primer used in the methods of the present
XX invention
SQ Sequence 36 BP; 9 A; 11 C; 10 G; 6 T; 0 U; 0 Other;
Query Match 26.2%; Score 17; DB 6; Length 36;
Best Local Similarity 100.0%; Pred. No. 27;
Matches 17; Conservative 0; Mismatches 0; Indels 0; Gaps 0;
QY 1 GATCCATGGAATCTGC 17
DB 2 GATCCATGGAATCTGC 18
RESULT 8
ABK47610
ID ABK47610 standard; DNA; 36 BP.
XX
XX ABK47610;
XX
XX 24-SEP-2002 (first entry)
XX
XX Interleukin-1 receptor antagonist (IL-1ra), PCR primer IRAS'.
XX
XX Human; elongation factor 1-alpha; retroviral vector; EF1a;
XX murine leukaemia virus vector; MLV; elongation factor 1 alpha;
XX gene therapy; interleukin-1 receptor antagonist; IL-1ra; PCR; primer; sr.
XX
XX Homo sapiens.
XX
XX WO200220810-A1.
XX
XX 14-MAR-2002.
XX
XX 08-SEP-2001; 2001WO-KR001515.
XX
XX 08-SEP-2000; 2000KR-00053613.
XX
XX (VIRO-) VIROMED LTD.
XX
XX Kim SY, Yu SS, Lee JT;
XX
XX WPI; 2002-065240/41.
XX
XX Retroviral vectors useful in gene therapy, containing no viral coding
XX sequences but which includes a human elongation factor 1 alpha non-coding
XX fragment.
XX
XX Example 5; Page 45; 47pp; English.
XX
XX The invention relates to a novel retroviral vector, derived from a murine
XX leukaemia virus (MLV) vector, which lacks viral coding sequences but
XX which includes: (i) part of the non-coding sequence (II) of elongation
XX factor 1 alpha (EF1a), as heterologous, gene-derived sequence, inserted
XX upstream of the multiple cloning site; and (ii) a mutation, downstream of
XX the splice acceptor, within (I). The retroviral vectors are useful as

CC gene therapy vectors. They are safe, i.e. they can not form replication-
CC competent retroviruses by homologous recombination; the heterologous
CC intron ensures efficient expression of foreign genes and the specified
CC viral titre is maintained. Also a heterologous promoter can be inserted
CC to increase viral titre in packaging lines derived from human cells. An
CC additional promoter (or internal ribosome entry site) can be provided to
CC allow expression of two or more genes. The present sequence represents a
CC PCR primer used to make the retroviral vector of the invention
XX
SQ Sequence 36 BP; 9 A; 11 C; 10 G; 6 T; 0 U; 0 Other;
Query Match 26.2%; Score 17; DB 6; Length 36;
Best Local Similarity 100.0%; Pred. No. 27;
Matches 17; Conservative 0; Mismatches 0; Indels 0; Gaps 0;
QY 1 GATCCATGGAATCTGC 17
DB 2 GATCCATGGAATCTGC 18
RESULT 9
AAH66339/C
ID AAH66339 standard; DNA; 990 BP.
XX
XX AAH66339;
XX
XX 26-SEP-2001 (first entry)
XX
XX C glutamicum coding sequence fragment SEQ ID NO: 1374.
XX
XX Corynebacterium; amino acid synthesis; vitamin; saccharide;
XX organic acid synthesis; ds.
XX
XX Corynebacterium glutamicum.
XX
XX EPI108790-A2.
XX
XX 20-JUN-2001.
XX
XX 18-DEC-2000; 2000EP-00127688.
XX
XX 16-DEC-1999; 99JP-00377484.
XX
XX 07-APR-2000; 2000JP-00159162.
XX
XX 03-AUG-2000; 2000JP-00280988.
XX
XX (KYOM) KYOMA HAKKO KOGYO KK.
XX
XX Nakagawa S, Mizoguchi H, Ando S, Hayashi M, Ochiai K, Yokoi H;
XX Tateishi N, Senoh A, Ikeda M, Ozaki A;
XX WPI; 2001-376931/40.
XX
XX P-PEDB; AAG91120.
XX
XX Novel polynucleotides derived from Corynebacterium bacteria, for identifying
XX expression point of a gene, measuring expression of a gene, analyzing
XX expression profile or pattern of a gene and identifying homologous gene.
XX
XX Claim 8; SEQ ID NO 1374; 246bp + Sequence Listing; English.
XX
XX The present invention provides a number of nucleotide and protein
XX sequences from the Corynebacterium bacteria Corynebacterium glutamicum. These
XX are useful for identifying the mutation point of a gene derived from a
XX mutant of corynebacterium bacteria, measuring expression amount and analysing
XX the expression profile or expression pattern of a gene derived from
XX Corynebacterium bacteria, and identifying a homologue of a gene derived from
XX corynebacterium bacteria. Corynebacterium bacteria are useful for producing amino
XX acids, nucleic acids, vitamins, saccharides and organic acids.
XX particularly L-lysine. The present sequence is a nucleic acid described
XX in the exemplification of the invention. Note: The sequence data for this
XX patent did not form part of the printed specification but was obtained
XX in electronic format directly from the European Patent Office

Sequence 990 BP; 196 A; 239 C; 310 G; 245 T; 0 U; 0 Other;
 Query Match 26.2%; Score 17; DB 5; Length 990;
 Best Local Similarity 100.0%; Pred. No. 26;
 Matches 17; Conservative 0; Mismatches 0; Indels 0; Gaps 0;

QY 1 GATCCATGGAATCTGC 17
 DB 688 GATCCATGGAATCTGC 672

RESULT 10
 AAF71853/c
 ID AAF71853 standard; DNA; 1113 BP.
 XX AAF71853;
 AC
 XX 30-APR-2001 (first entry)
 DT
 XX Corynebacterium glutamicum MP protein nucleotide sequence SEQ ID NO:201.
 DE
 XX Corynebacterium glutamicum; metabolic pathway protein; MP protein;
 KW fine chemical production; microorganism; organic acid; nucleoside;
 KW nonproteinogenic amino acid; purine base; pyrimidine base; nucleotide;
 KW lipid; saturated fatty acid; unsaturated fatty acid; diol; vitamin;
 KW carbohydrate; aromatic compound; cofactor; polyketide; enzyme; ds.
 XX
 OS Corynebacterium glutamicum.
 XX
 XX W0200100843-A2.
 XX
 PD 04-JAN-2001.
 XX
 PF 23-JUN-2000; 2000MO-IB000923.
 XX
 PR 25-JUN-1999; 99US-0141031P
 PR 01-JUL-1999; 99DE-01030676.
 PR 02-JUL-1999; 99US-0142101P.
 PR 08-JUL-1999; 99DE-01031415.
 PR 08-JUL-1999; 99DE-01031418.
 PR 08-JUL-1999; 99DE-01031419.
 PR 08-JUL-1999; 99DE-01031420.
 PR 08-JUL-1999; 99DE-01031424.
 PR 08-JUL-1999; 99DE-01031428.
 PR 08-JUL-1999; 99DE-01031434.
 PR 08-JUL-1999; 99DE-01031435.
 PR 08-JUL-1999; 99DE-01031443.
 PR 08-JUL-1999; 99DE-01031453.
 PR 08-JUL-1999; 99DE-01031457.
 PR 08-JUL-1999; 99DE-01031465.
 PR 08-JUL-1999; 99DE-01031478.
 PR 08-JUL-1999; 99DE-01031510.
 PR 08-JUL-1999; 99DE-01031541.
 PR 08-JUL-1999; 99DE-01031573.
 PR 08-JUL-1999; 99DE-01031592.
 PR 08-JUL-1999; 99DE-01031632.
 PR 08-JUL-1999; 99DE-01031634.
 PR 08-JUL-1999; 99DE-01031636.
 PR 09-JUL-1999; 99DE-01032125.
 PR 09-JUL-1999; 99DE-01032126.
 PR 09-JUL-1999; 99DE-01032130.
 PR 09-JUL-1999; 99DE-01032186.
 PR 09-JUL-1999; 99DE-01032206.
 PR 09-JUL-1999; 99DE-01032227.
 PR 09-JUL-1999; 99DE-01032228.
 PR 09-JUL-1999; 99DE-01032229.
 PR 09-JUL-1999; 99DE-01032230.
 PR 14-JUL-1999; 99DE-01032822.
 PR 14-JUL-1999; 99DE-01032826.
 PR 14-JUL-1999; 99DE-01032828.
 PR 14-JUL-1999; 99DE-01033004.
 PR 14-JUL-1999; 99DE-01033005.
 PR 14-JUL-1999; 99DE-01033006.

PR 12-AUG-1999; 99US-0148613P.
 PR 27-AUG-1999; 99DE-01040764.
 PR 27-AUG-1999; 99DE-01040765.
 PR 27-AUG-1999; 99DE-01040766.
 PR 31-AUG-1999; 99DE-01040832.
 PR 31-AUG-1999; 99DE-01041378.
 PR 31-AUG-1999; 99DE-01041379.
 PR 31-AUG-1999; 99DE-01041380.
 PR 31-AUG-1999; 99DE-01041394.
 PR 31-AUG-1999; 99DE-01041396.
 PR 03-SEP-1999; 99DE-01042076.
 PR 03-SEP-1999; 99DE-01042079.
 PR 03-SEP-1999; 99DE-01042086.
 PR 03-SEP-1999; 99DE-01042087.
 PR 03-SEP-1999; 99DE-01042088.
 PR 03-SEP-1999; 99DE-01042095.
 PR 03-SEP-1999; 99DE-01042124.
 PR 03-SEP-1999; 99DE-01042129.
 PR 09-MAR-2000; 2000US-0187970P.
 XX
 PA (BAD) BASF AG.
 XX
 XX Pompejus M, Krosger B, Schroeder H, Zelder O, Haberhauer G;
 XX
 DR MPI; 2001-137957/14.
 DR P-PSDB; AAB79734.
 XX
 PT Nucleic acids from Corynebacterium glutamicum encoding metabolic pathway
 PT proteins, useful for producing fine chemicals in microorganisms,
 PT including organic acids, nonproteinogenic amino acids, and purine and
 PT pyrimidine bases.
 XX
 PS Claim 3; Page 446-447; 1737pp; English.
 XX
 CC AAF71753 to AAF72330 encode the Corynebacterium glutamicum metabolic
 CC pathway (MP) proteins given in AAB79634 to AAB80211. The C. glutamicum MP
 CC nucleic acids are useful for the production of fine chemicals in
 CC microorganisms, including organic acids, nonproteinogenic amino acids,
 CC purine and pyrimidine bases, nucleosides, nucleotides, lipids, saturated
 CC and unsaturated fatty acids, diols, carbohydrates, aromatic compounds,
 CC vitamins, cofactors, polyketides and enzymes
 XX
 SQ Sequence 1113 BP; 220 A; 271 C; 348 G; 274 T; 0 U; 0 Other;
 Query Match 26.2%; Score 17; DB 4; Length 1113;
 Best Local Similarity 100.0%; Pred. No. 26;
 Matches 17; Conservative 0; Mismatches 0; Indels 0; Gaps 0;

QY 1 GATCCATGGAATCTGC 17
 DB 788 GATCCATGGAATCTGC 772

RESULT 11
 AAS96115/c
 ID AAS96115 standard; DNA; 1113 BP.
 XX AAS96115;
 AC
 XX 26-FEB-2002 (first entry)
 DT
 XX C. glutamicum gene #40 encoding metabolic pathway protein.
 DE
 XX Metabolic pathway protein; MP; lysine biosynthesis pathway;
 KW methionine biosynthesis pathway; large-scale production of fine chemical;
 KW Corynebacterium diptheriae; diptheria; ds.
 XX
 OS Corynebacterium glutamicum.
 XX
 XX W0200106573-A2.
 XX
 PN 13-SEP-2001.
 XX

XX 22-DEC-2000; 2000WO-IB002035.
 PF 09-MAR-2000; 2000US-0187970P.
 PR 23-JUN-2000; 2000US-00606740.
 XX (BADI) BASF AG.
 PA
 PI Pompejus M, Kroegeer B, Schroeder H, Zelder O, Habernauer G;
 PI Kim U, Lee H, Hwang B;
 XX WPI; 2001-582839/65.
 DR P-PSDB; AAU71905.
 XX Nucleic acids encoding metabolic pathway proteins from Corynebacterium
 PT glutamicum, useful for producing methionine and lysine in Corynebacterium
 PT and Brevibacterium.
 XX
 PS Disclosure; page 256-257; 316pp; English.
 XX The present invention relates to the isolation of novel Corynebacterium
 CC glutamicum genes encoding metabolic pathway (MP) proteins (AAU71863-
 CC AAU71922). The metabolic pathway proteins of the invention include
 CC enzymes involved in the lysine and methionine biosynthetic pathways. The
 CC polynucleotide sequences of the invention can be used for the large-scale
 CC production and/or modulation of expression of fine chemicals such as
 CC lysine and methionine. The sequences of the invention may be used to
 CC identify C. glutamicum and related organisms e.g. C. diphtheriae in a
 CC subject to detect diphtheria. AAS96073-AAS96132 represent C. glutamicum
 CC genes encoding the novel metabolic pathway proteins of the invention
 XX
 SQ Sequence 113 BP; 220 A; 271 C; 348 G; 274 T; 0 U; 0 Other;
 Query Match 26.2%; Score 17; DB 4; Length 113;
 Best Local Similarity 100.0%; Pred. No. 26;
 Matches 17; Conservative 0; Mismatches 0; Indels 0; Gaps 0;
 QY 1 GATCCATGGAATCTGC 17
 Db 788 GATCCATGGAATCTGC 772
 RESULT 12
 ID AAS74960 standard; CDNA; 2055 BP.
 AC AAS74960;
 XX
 DT 13-FEB-2002 (first entry)
 DE DNA encoding novel human diagnostic protein #10764.
 XX
 KW Human; chromosome mapping; gene mapping; gene therapy; forensic;
 KW food supplement; medical imaging; diagnostic; genetic disorder; ss.
 XX
 OS Homo sapiens.
 XX
 PN MO200175067-A2.
 PD 11-OCT-2001.
 PF 30-MAR-2001; 2001WO-US008631.
 PR 31-MAR-2000; 2000US-00540217.
 PR 23-AUG-2000; 2000US-00649167.
 XX
 PA (HYSE-) HYSEQ INC.
 XX
 PI Dmanac RT, Liu C, Tang YT;
 XX
 DR WPI; 2001-639362/73.
 DR P-PSDB; ABG10773.
 XX

PT New isolated polynucleotide and encoded polypeptides, useful in
 PT diagnostics, forensics, gene mapping, identification of mutations
 PT responsible for genetic disorders or other traits and to assess
 PT biodiversity.
 XX
 XX Claim 1; SEQ ID NO 10764; 103pp; English.
 XX
 CC The invention relates to isolated polynucleotide (I) and polypeptide (II)
 CC sequences. (I) is useful as hybridisation probes, polymerase chain
 CC reaction (PCR) primers, oligomers, and for chromosome and gene mapping,
 CC and in recombinant production or (II). The polynucleotides are also used
 CC in diagnostics as expressed sequence tags for identifying expressed
 CC genes. (I) is useful in gene therapy techniques to restore normal
 CC activity of (II) or to treat disease states involving (II). (II) is
 CC useful for generating antibodies against it, detecting or quantitating a
 CC polypeptide in tissue, as molecular weight markers and as a food
 CC supplement. (II) and its binding partners are useful in medical imaging
 CC of sites expressing (II). (I) and (II) are useful for treating disorders
 CC involving aberrant protein expression or biological activity. The
 CC polypeptide and polynucleotide sequences have applications in
 CC diagnostics, forensics, gene mapping, identification of mutations
 CC responsible for genetic disorders or other traits to assess biodiversity
 CC and to produce other types of data and products dependent on DNA and
 CC amino acid sequences. AAS64197-AAS94564 represent novel human diagnostic
 CC coding sequences of the invention. Note: The sequence data for this
 CC patent did not appear in the printed specification, but was obtained in
 CC electronic format directly from WIPO at
 CC ftp.wipo.int/pub/published_pct_sequences
 XX
 SQ Sequence 2055 BP; 469 A; 581 C; 604 G; 401 T; 0 U; 0 Other;
 Query Match 26.2%; Score 17; DB 5; Length 2055;
 Best Local Similarity 100.0%; Pred. No. 26;
 Matches 17; Conservative 0; Mismatches 0; Indels 0; Gaps 0;
 QY 40 AAACCTGCTGATGAGC 56
 Db 98 AAACCTGCTGATGAGC 114
 RESULT 13
 ID ACA43988 standard; DNA; 2877 BP.
 AC ACA43988;
 XX
 DT 19-JUN-2003 (first entry)
 DE Prokaryotic essential gene #25645.
 XX
 KW Antisense; ds; prokaryotic essential gene; cell proliferation;
 KW drug design; gene.
 XX
 OS Pseudomonas putida.
 XX
 PN WO200277183-A2.
 PD 03-OCT-2002.
 PF 21-MAR-2002; 2002WO-US009107.
 PR 21-MAR-2001; 2001US-00815242.
 PR 06-SEP-2001; 2001US-00948993.
 PR 25-OCT-2001; 2001US-0342923P.
 PR 08-FEB-2002; 2002US-00072851.
 PR 06-MAR-2002; 2002US-0362699P.
 XX
 PA (ELIT-) ELITRA PHARM INC.
 XX
 PI Wang J, Zamudio C, Malone C, Haselbeck R, Ohlsen KL, Zyskind JW;
 PI Wall D, Trawick JD, Carr GT, Yamamoto R, Foreyth RA, Xu HH;
 XX
 DR WPI; 2003-029926/02.
 XX

DR P-PSDB; AEU40118.

XX New antisense nucleic acids, useful for identifying proteins or screening
PT for homologous nucleic acids, required for cellular proliferation to
PT isolate candidate molecules for rational drug discovery programs.

PS Claim 14; SEQ ID NO 31858; 1766pp; English.

XX The invention relates to an isolated nucleic acid comprising any one of
CC the 6213 antisense sequences given in the specification where expression
CC of the nucleic acid inhibits proliferation of a cell. Also included are:
CC (1) a vector comprising a promoter operably linked to the nucleic acid
CC encoding a polypeptide whose expression is inhibited by the antisense
CC nucleic acid; (2) a host cell containing the vector; (3) an isolated
CC polypeptide or its fragment whose expression is inhibited by the
CC antisense nucleic acid; (4) an antibody capable of specifically binding
CC the polypeptide; (5) producing the polypeptide; (6) inhibiting cellular
CC proliferation or the activity of a gene in an operon required for
CC proliferation; (7) identifying a compound that influences the activity of
CC the gene product or that has an activity against a biological pathway
CC required for proliferation, or that inhibits cellular proliferation; (8)
CC identifying a gene required for cellular proliferation or the biological
CC pathway in which a proliferation-regulated gene or its gene product lies
CC or a gene on which the test compound that inhibits proliferation of an
CC organism acts; (9) manufacturing an antibiotic; (10) profiling a
CC compound's activity; (11) a culture comprising strains in which the gene
CC product is overexpressed or underexpressed; (12) determining the extent
CC to which each of the strains is present in a culture or collection of
CC strains; or (13) identifying the target of a compound that inhibits the
CC proliferation of an organism. The antisense nucleic acids are useful for
CC identifying proteins or screening for homologous nucleic acids required
CC for cellular proliferation to isolate candidate molecules for rational
CC drug discovery programs, or for screening homologous nucleic acids
CC required for proliferation in cells other than *S. aureus*, *S. typhimurium*,
CC *K. pneumoniae* or *P. aeruginosa*. The present sequence is one of the target
CC prokaryotic essential genes. Note: The sequence data for this patent did
CC not form part of the printed specification, but was obtained in
CC electronic format directly from WIPO at
CC [ftp://wipo.int/pub/published_pct_sequences](http://wipo.int/pub/published_pct_sequences)

SQ Sequence 2877 BP; 629 A; 981 C; 802 G; 465 T; 0 U; 0 Other;

Query Match 26.2%; Score 17; DB 8; Length 2877;

Best Local Similarity 100.0%; Pred. No. 26; Mismatches 0; Indels 0; Gaps 0;

Matches 17; Conservative 0; Mismatches 0; Indels 0; Gaps 0;

QY 44 CCTGCTGATGACACCC 60

DB 675 CCTGCTGATGACACCC 691

RESULT 14

AAS54085 standard; DNA; 2892 BP.

AC AAS54085;

DT 13-FEB-2002 (first entry)

DE Pseudomonas aeruginosa DNA for cellular proliferation protein #216.

XX Antisense; ds; prokaryotic cellular proliferation gene; antibiotic;
KM antibacterial; drug design.

XX Pseudomonas aeruginosa.

XX WO200170955-A2.

XX 27-SEP-2001.

XX 21-MAR-2001; 2001WO-US009180.

XX 21-MAR-2000; 2000US-0191078P.

PR 23-MAY-2000; 2000US-0206848P.

PR 26-MAY-2000; 2000US-0207227P.

PR 27-OCT-2000; 2000US-0242578P.

PR 27-NOV-2000; 2000US-0253625P.

PR 22-DEC-2000; 2000US-0257531P.

PR 16-FEB-2001; 2001US-0269308P.

XX (ELIT-) ELITRA PHARM INC.

XX Haselbeck R, Ohlsen KL, Zyskind JW, Wall D, Trawick JD, Carr CJ;

PI Yamamoto RT, Xu HH;

XX WPI, 2001-611495/70.

DR P-PSDB; AEU40118.

XX New polynucleotides for the identification and development of

PT antibiotics, comprise sequences of antisense nucleic acids.

XX Claim 27; SEQ ID NO 7722; 511pp; English.

XX The invention relates to antisense inhibitors of genes essential to
CC prokaryotic cellular proliferation, their use in identifying the genes,
CC their use in the discovery of novel antibiotics, the essential genes
CC themselves and the encoded proteins. The prokaryotes used are *Escherichia*
CC coli, *Staphylococcus aureus*, *Salmonella typhi*, *Klebsiella pneumoniae*,
CC *Pseudomonas aeruginosa* and *Enterococcus faecalis*. The invention is also
CC useful for the identification of potential new targets for antibiotic
CC development. The antisense nucleic acids can also be used to identify
CC proteins used in proliferation, to express these proteins, and to obtain
CC antibodies capable of binding to the expressed proteins. The proteins can
CC be used to screen compounds in rational drug discovery programmes. The
CC antisense nucleic acid sequence is also useful to screen for homologous
CC nucleic acids which are required for cell proliferation in a wide variety
CC of organisms. The present sequence encodes an essential prokaryotic
CC cellular proliferation protein. Note: The sequence data for this patent
CC did not form part of the printed specification, but was obtained in
CC electronic format directly from WIPO at
CC [ftp://wipo.int/pub/published_pct_sequences](http://wipo.int/pub/published_pct_sequences)

SQ Sequence 2892 BP; 594 A; 1063 C; 824 G; 411 T; 0 U; 0 Other;

Query Match 26.2%; Score 17; DB 4; Length 2892;

Best Local Similarity 100.0%; Pred. No. 26; Mismatches 0; Indels 0; Gaps 0;

Matches 17; Conservative 0; Mismatches 0; Indels 0; Gaps 0;

QY 44 CCTGCTGATGACACCC 60

DB 675 CCTGCTGATGACACCC 691

RESULT 15

ACA42199 standard; DNA; 2892 BP.

AC ACA42199;

DT 19-JUN-2003 (first entry)

DE Prokaryotic essential gene #23856.

XX Antisense; ds; prokaryotic essential gene; cell proliferation;
KM drug design; gene.

XX Pseudomonas aeruginosa.

XX WO200277183-A2.

XX 03-OCT-2002.

XX 21-MAR-2002; 2002WO-US009107.

XX 21-MAR-2001; 2001US-00815242.

XX 06-SEP-2001; 2001US-00948993.

PR 25-OCT-2001; 2001US-0342923P.
 PR 08-FEB-2002; 2002US-0007285P.
 PR 06-MAR-2002; 2002US-0362699P.
 XX
 XX (ELIT-) ELITRA PHARM INC.
 XX
 PI Wang L, Zamudio C, Malone C, Haselbeck R, Ohlsen KL, Zyskind JW;
 PI Wall D, Trawick JD, Carr GT, Yamamoto R, Forsyth RA, Xu HH;
 XX
 DR WPI; 2003-029926/02;
 DR P-PSDB; ABU38329.
 XX
 PT New antisense nucleic acids, useful for identifying proteins or screening
 PT for homologous nucleic acids required for cellular proliferation to
 PT isolate candidate molecules for rational drug discovery programs.
 XX
 PS Claim 14; SEQ ID NO 30069; 1766bp; English.
 XX
 CC The invention relates to an isolated nucleic acid comprising any one of
 CC the 6213 antisense sequences given in the specification where expression
 CC of the nucleic acid inhibits proliferation of a cell. Also included are:
 CC (1) a vector comprising a promoter operably linked to the nucleic acid
 CC encoding a polypeptide whose expression is inhibited by the antisense
 CC nucleic acid; (2) a host cell containing the vector; (3) an isolated
 CC polypeptide or its fragment whose expression is inhibited by the
 CC antisense nucleic acid; (4) an antibody capable of specifically binding
 CC the polypeptide; (5) producing the polypeptide; (6) inhibiting cellular
 CC proliferation or the activity of a gene in an operon required for
 CC proliferation; (7) identifying a compound that influences the activity of
 CC the gene product or that has an activity against a biological pathway
 CC required for proliferation, or that inhibits cellular proliferation; (8)
 CC identifying a gene required for cellular proliferation or the biological
 CC pathway in which a proliferation-required gene or its gene product lies
 CC or a gene on which the test compound that inhibits proliferation of an
 CC organism acts; (9) manufacturing an antibiotic; (10) profiling a
 CC compound's activity; (11) a culture comprising strains in which the gene
 CC product is overexpressed or underexpressed; (12) determining the extent
 CC to which each of the strains is present in a culture or collection of
 CC strains; or (13) identifying the target of a compound that inhibits the
 CC proliferation of an organism. The antisense nucleic acids are useful for
 CC identifying proteins or screening for homologous nucleic acids required
 CC for cellular proliferation to isolate candidate molecules for rational
 CC drug discovery programs, or for screening homologous nucleic acids
 CC required for proliferation in cells other than *S. aureus*, *S. typhimurium*,
 CC *K. pneumoniae* or *P. aeruginosa*. The present sequence is one of the target
 CC prokaryotic essential genes. Note: The sequence data for this patent did
 CC not form part of the printed specification, but was obtained in
 CC electronic format directly from WIPO at
 CC ftp.wipo.int/pub/published_pct_sequences
 XX
 SQ Sequence 2892 BP; 594 A; 1063 C; 824 G; 411 T; 0 U; 0 Other;
 XX
 Query Match 26.2%; Score 17; DB 8; Length 2892;
 Best Local Similarity 100.0%; Pred. No. 26;
 Matches 17; Conservative 0; Mismatches 0; Indels 0; Gaps 0;
 QY 44 CCTGCTGATGACACCC 60
 DB 675 CCTGCTGATGACACCC 691

Search completed: December 22, 2004, 22:44:07
 Job time : 234.662 secs

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RESULT 2
US-08-864-357F-13/c
Sequence 13, Application US/08864357F
Patent No. 6255281
GENERAL INFORMATION:
APPLICANT: Ciaragen, Inc. & NIH
TITLE OF INVENTION: Use of Recombinant Human Uteroglobulin in Treatment of Inflammation
TITLE OF INVENTION: Fibrotic Conditions
FILE REFERENCE: 1.6142/2
CURRENT APPLICATION NUMBER: US/08/864,357F
CURRENT FILING DATE: 1997-05-28
NUMBER OF SEQ. ID NOS.: 22
SOFTWARE: PatentIn version 3.0
SEQ ID NO 13
LENGTH: 42
TYPE: DNA

```

ORGANISM: artificial
FEATURE:
OTHER INFORMATION: primer sequence
US-08-864-357F-13

Query Match 64.6%; Score 42; DB 3; Length 42;
Best Local Similarity 100.0%; Pred. No. 5,6e-14;
Matches 42; Conservative 0; Mismatches 0; Indels 0; Gaps 0;

Qy 1 GATCCATGAAGAATCGCCGCTCTTCCAGCGCTTATCATAA 42
Db 42 GATCCATGAAGAATCGCCGCTCTTCCAGCGCTTATCATAA 1

RESULT 3
US-08-864-357F-12/c
Sequence 12, Application US/08864357F
Patent No. 6255281
GENERAL INFORMATION:
APPLICANT: Clargen, Inc. & NIH
TITLE OF INVENTION: Use of Recombinant Human Uteroglobulin in Treatment of Inflammato
FILE REFERENCE: 116142/2
CURRENT APPLICATION NUMBER: US/08/864,357F
CURRENT FILING DATE: 1997-05-28
NUMBER OF SEQ ID NOS: 22
SOFTWARE: Patent version 3.0
SEQ ID NO 12
LENGTH: 60
TYPE: DNA
ORGANISM: artificial
FEATURE:
OTHER INFORMATION: primer sequence
US-08-864-357F-12

Query Match 35.4%; Score 23; DB 3; Length 60;
Best Local Similarity 100.0%; Pred. No. 0.0016;
Matches 23; Conservative 0; Mismatches 0; Indels 0; Gaps 0;

Qy 43 CCTGCTGATGACACCCCGCTCC 65
Db 60 CCTGCTGATGACACCCCGCTCC 38

RESULT 4
US-09-252-991A-2879
Sequence 2879, Application US/09252991A
Patent No. 6551795
GENERAL INFORMATION:
APPLICANT: Marc J. Rubenfield et al.
TITLE OF INVENTION: NUCLEIC ACID AND AMINO ACID SEQUENCES RELATING TO PSEUDOMONAS
FILE REFERENCE: 107196.136
CURRENT APPLICATION NUMBER: US/09/252,991A
CURRENT FILING DATE: 1999-02-18
PRIOR APPLICATION NUMBER: US 60/074,788
PRIOR FILING DATE: 1998-02-18
PRIOR APPLICATION NUMBER: US 60/094,190
PRIOR FILING DATE: 1998-07-27
NUMBER OF SEQ ID NOS: 33142
SEQ ID NO 2879
LENGTH: 2997
TYPE: DNA
ORGANISM: Pseudomonas aeruginosa
US-09-252-991A-2879

Query Match 26.2%; Score 17; DB 4; Length 2997;
Best Local Similarity 100.0%; Pred. No. 3.1;
Matches 17; Conservative 0; Mismatches 0; Indels 0; Gaps 0;

Qy 44 CCTGCTGATGACACCC 60
Db 780 CCTGCTGATGACACCC 756

RESULT 5
US-09-252-991A-3091/c
Sequence 3091, Application US/09252991A
Patent No. 6551795
GENERAL INFORMATION:
APPLICANT: Marc J. Rubenfield et al.
TITLE OF INVENTION: NUCLEIC ACID AND AMINO ACID SEQUENCES RELATING TO PSEUDOMONAS
FILE REFERENCE: 107196.136
CURRENT APPLICATION NUMBER: US/09/252,991A
CURRENT FILING DATE: 1999-02-18
PRIOR APPLICATION NUMBER: US 60/074,788
PRIOR FILING DATE: 1998-02-18
PRIOR APPLICATION NUMBER: US 60/094,190
PRIOR FILING DATE: 1998-07-27
NUMBER OF SEQ ID NOS: 33142
SEQ ID NO 3091
LENGTH: 3009
TYPE: DNA
ORGANISM: Pseudomonas aeruginosa
US-09-252-991A-3091

Query Match 26.2%; Score 17; DB 4; Length 3009;
Best Local Similarity 100.0%; Pred. No. 3.1;
Matches 17; Conservative 0; Mismatches 0; Indels 0; Gaps 0;

Qy 44 CCTGCTGATGACACCC 60
Db 2428 CCTGCTGATGACACCC 2412

RESULT 6
US-09-489-039A-1155
Sequence 1155, Application US/09489039A
Patent No. 6610836
GENERAL INFORMATION:
APPLICANT: Gary Breton et. al
TITLE OF INVENTION: NUCLEIC ACID AND AMINO ACID SEQUENCES RELATING TO KLEBSIELLA
FILE REFERENCE: 2709.2004001
CURRENT APPLICATION NUMBER: US/09/489,039A
CURRENT FILING DATE: 2000-01-27
PRIOR APPLICATION NUMBER: US 60/117,747
PRIOR FILING DATE: 1999-01-29
NUMBER OF SEQ ID NOS: 14342
SEQ ID NO 1155
LENGTH: 894
TYPE: DNA
ORGANISM: Klebsiella pneumoniae
US-09-489-039A-1155

Query Match 24.6%; Score 16; DB 4; Length 894;
Best Local Similarity 100.0%; Pred. No. 11;
Matches 16; Conservative 0; Mismatches 0; Indels 0; Gaps 0;

Qy 40 AAACCTGCTGATGA 55
Db 527 AAACCTGCTGATGA 542

RESULT 7
US-09-489-039A-3704
Sequence 3704, Application US/09489039A
Patent No. 6610836
GENERAL INFORMATION:
APPLICANT: Gary Breton et. al
TITLE OF INVENTION: NUCLEIC ACID AND AMINO ACID SEQUENCES RELATING TO KLEBSIELLA
FILE REFERENCE: 2709.2004001
CURRENT APPLICATION NUMBER: US/09/489,039A
CURRENT FILING DATE: 2000-01-27

PRIOR APPLICATION NUMBER: US 60/117,747
 PRIOR FILING DATE: 1999-01-29
 NUMBER OF SEQ ID NOS: 14342
 SEQ ID NO 3704
 LENGTH: 1119
 TYPE: DNA
 ORGANISM: Klebsiella pneumoniae
 US-09-489-039A-3704

Query Match 24.6% Score 16; DB 4; Length 1119;
 Best Local Similarity 100.0%; Pred. No. 11;
 Matches 16; Conservative 0; Mismatches 0; Indels 0; Gaps 0;

QY 45 CTGCTGATGACACCC 60
 DB 937 CTGCTGATGACACCC 952

RESULT 8
 US-08-858-207A-101
 Sequence 101, Application US/08858207A
 Patent No. 6348328

GENERAL INFORMATION:
 APPLICANT: Black, Michael
 APPLICANT: Hodgson, John
 APPLICANT: Knowles, David
 APPLICANT: Nicholas, Richard
 APPLICANT: Stedola, Robert
 TITLE OF INVENTION: No. 6348328e1 Compounds
 NUMBER OF SEQUENCES: 552
 CORRESPONDENCE ADDRESS:
 ADDRESSEE: SmithKline Beecham Corporation
 STREET: 709 Swedeland Road
 CITY: King of Prussia
 STATE: PA
 COUNTRY: USA
 ZIP: 19406-0939

COMPUTER READABLE FORM:
 MEDIUM TYPE: Diskette
 COMPUTER: IBM Compatible
 OPERATING SYSTEM: DOS
 SOFTWARE: FastSeq for Windows Version 2.0
 CURRENT APPLICATION DATA:
 APPLICATION NUMBER: US/08/858,207A
 FILING DATE: 09-MAY-1997
 CLASSIFICATION: 435
 PRIOR APPLICATION DATA:
 APPLICATION NUMBER: 60/017670
 FILING DATE: 14-MAY-1996

ATTORNEY/AGENT INFORMATION:
 NAME: Gimm, Edward R
 REGISTRATION NUMBER: 38,891
 REFERENCE/DOCKET NUMBER: P50475
 TELECOMMUNICATION INFORMATION:
 TELEPHONE: 610-270-4478
 TELEFAX: 610-270-5090
 TELEX:

INFORMATION FOR SEQ ID NO: 101:
 SEQUENCE CHARACTERISTICS:
 LENGTH: 1155 base pairs
 TYPE: nucleic acid
 STRANDEDNESS: single
 TOPOLOGY: linear
 US-08-858-207A-101

Query Match 24.6% Score 16; DB 3; Length 1155;
 Best Local Similarity 100.0%; Pred. No. 11;
 Matches 16; Conservative 0; Mismatches 0; Indels 0; Gaps 0;

QY 3 TCCTAGGAATCTGCC 18
 DB 1011 TCCTAGGAATCTGCC 1026

RESULT 9
 US-09-023-655-481/C
 Sequence 481, Application US/09023655
 Patent No. 6607879

GENERAL INFORMATION:
 APPLICANT: Cocks, Benjamin G.
 APPLICANT: Susan G. Stuart
 APPLICANT: Jeffrey J. Seilhamer
 TITLE OF INVENTION: COMPOSITION FOR THE DETECTION OF BLOOD CELL GENE
 TITLE OF INVENTION: EXPRESSION
 NUMBER OF SEQUENCES: 1508
 CORRESPONDENCE ADDRESS:
 ADDRESSEE: INCYTE PHARMACEUTICALS, INC.
 STREET: 3174 PORTER DRIVE
 CITY: PALO ALTO
 STATE: CALIFORNIA
 COUNTRY: USA
 ZIP: 94304

COMPUTER READABLE FORM:
 MEDIUM TYPE: Floppy disk
 COMPUTER: IBM PC compatible
 OPERATING SYSTEM: PC-DOS/MS-DOS
 SOFTWARE: Word Perfect 6.1 for Windows/MS-DOS 6.2
 CURRENT APPLICATION DATA:
 APPLICATION NUMBER: US/09/023,655
 FILING DATE: HEREMITH
 CLASSIFICATION:
 PRIOR APPLICATION DATA:
 APPLICATION NUMBER:
 FILING DATE:

ATTORNEY/AGENT INFORMATION:
 NAME: Zeller, Karen J
 REGISTRATION NUMBER: 37,071
 REFERENCE/DOCKET NUMBER: PA-0001 US
 TELECOMMUNICATION INFORMATION:
 TELEPHONE: (650) 845-4166
 TELEFAX: (650) 855-0555
 INFORMATION FOR SEQ ID NO: 481:
 SEQUENCE CHARACTERISTICS:
 LENGTH: 1434 base pairs
 TYPE: nucleic acid
 STRANDEDNESS: single
 TOPOLOGY: linear
 IMMEDIATE SOURCE:
 LIBRARY: CONNUT01
 CLONE: 1908804
 US-09-023-655-481

Query Match 24.6% Score 16; DB 4; Length 1434;
 Best Local Similarity 100.0%; Pred. No. 11;
 Matches 16; Conservative 0; Mismatches 0; Indels 0; Gaps 0;

QY 14 CTGCCCGCTTCCAG 29
 DB 26 CTGCCCGCTTCCAG 11

RESULT 10
 US-09-566-921-61/C
 Sequence 61, Application US/09566921
 Patent No. 6682888

GENERAL INFORMATION:
 APPLICANT: Loring, Jeanne F.
 APPLICANT: Tingley, Debora W.
 APPLICANT: Edwards, Carla M.
 TITLE OF INVENTION: GENES EXPRESSED IN ALZHEIMER'S DISEASE
 FILE REFERENCE: PA-0024 US
 CURRENT APPLICATION NUMBER: US/09/566,921
 CURRENT FILING DATE: 2000-05-05
 NUMBER OF SEQ ID NOS: 138
 SOFTWARE: PERL Program

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; SEQ ID NO 61
; LENGTH: 3911
; TYPE: DNA
; ORGANISM: Homo sapiens
; FEATURE:
; NAME/KEY: misc_feature
; OTHER INFORMATION: Incyte ID No. 6682888 166400.9
US-09-566-921-61

```

```

Query Match      24.6%; Score 16; DB 4; Length 3911;
Best Local Similarity 100.0%; Pred. No. 11;
Matches 16; Conservative 0; Mismatches 0; Indels 0; Gaps 0;

```

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Qy      14  CTGCCCGCTTTCCAG 29
Db      2494 CTGCCCGCTTTCCAG 2479

```

```

RESULT 11
US-09-166-350-32
; Sequence 32, Application US/09166350A
; Patent No. 6440663
; GENERAL INFORMATION:
; APPLICANT: Scanlan, Matthew
; APPLICANT: Chen, Yao
; APPLICANT: Stockert, Elisabeth
; APPLICANT: Old, Lloyd
; APPLICANT: Uager, Elke
; APPLICANT: Knuth, Alex
; TITLE OF INVENTION: Renal Cancer Associated Antigens and
; FILE REFERENCE: L0461/7051
; CURRENT APPLICATION NUMBER: US/09/166,350A
; EARLIER FILING DATE: 1998-10-05
; EARLIER APPLICATION NUMBER: US 09/166,350
; EARLIER FILING DATE: 1998-10-05
; NUMBER OF SEQ ID NOS: 35
; SOFTWARE: FastSeq for Windows Version 3.0
; SEQ ID NO 32
; LENGTH: 4169
; TYPE: DNA
; ORGANISM: Homo sapiens
US-09-166-350-32

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Query Match      24.6%; Score 16; DB 4; Length 4169;
Best Local Similarity 100.0%; Pred. No. 11;
Matches 16; Conservative 0; Mismatches 0; Indels 0; Gaps 0;

```

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Qy      43  CCTCTGATGACAC 58
Db      3093 CCTCTGATGACAC 3108

```

```

RESULT 12
US-09-513-999C-11687/c
; Sequence 11687, Application US/09513999C
; Patent No. 6783961
; GENERAL INFORMATION:
; APPLICANT: Dumas Milne Edwards, J.B.
; APPLICANT: Duclert, A.Y.
; TITLE OF INVENTION: Expressed Sequence Tags and Encoded Human Proteins.
; Patent No. 6783961
; FILE REFERENCE: 59, US2, REG
; CURRENT APPLICATION NUMBER: US/09/513,999C
; CURRENT FILING DATE: 2000-02-24
; PRIOR APPLICATION NUMBER: US 60/122,487
; PRIOR FILING DATE: 1999-02-26
; NUMBER OF SEQ ID NOS: 36681
; SOFTWARE: Patent.pm
; SEQ ID NO 11687
; LENGTH: 301
; TYPE: DNA

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; ORGANISM: Homo sapiens
; FEATURE:
; NAME/KEY: misc_feature
; LOCATION: 244
; OTHER INFORMATION: n=a, g, c or t
; FEATURE:
; NAME/KEY: misc_feature
; LOCATION: 274
; OTHER INFORMATION: r=a or g
US-09-513-999C-11687

```

```

Query Match      23.1%; Score 15; DB 4; Length 301;
Best Local Similarity 100.0%; Pred. No. 39;
Matches 15; Conservative 0; Mismatches 0; Indels 0; Gaps 0;

```

```

Qy      17  CCGCTTTCCAGCG 31
Db      32  CCGCTTTCCAGCG 18

```

```

RESULT 13
US-09-702-705-1590
; Sequence 1590, Application US/09702705
; Patent No. 6504010
; GENERAL INFORMATION:
; APPLICANT: Wang, Tongtong
; APPLICANT: Bangur, Chaitanya S.
; APPLICANT: Lodes, Michael A.
; APPLICANT: Fanger, Gary
; APPLICANT: Vedvick, Tom
; APPLICANT: Carter, Darrick
; APPLICANT: Retter, Marc
; APPLICANT: Mannion, Jane
; TITLE OF INVENTION: COMPOSITIONS AND METHODS FOR THE THERAPY AND
; FILE REFERENCE: 210121.4/78C14
; CURRENT APPLICATION NUMBER: US/09/702,705
; CURRENT FILING DATE: 2000-10-30
; NUMBER OF SEQ ID NOS: 1833
; SOFTWARE: FastSeq for Windows Version 3.0
; SEQ ID NO 1590
; LENGTH: 434
; TYPE: DNA
; ORGANISM: Homo sapiens
; FEATURE:
; NAME/KEY: misc_feature
; LOCATION: (1)...(434)
; OTHER INFORMATION: n = A,T,C or G
US-09-702-705-1590

```

```

Query Match      23.1%; Score 15; DB 4; Length 434;
Best Local Similarity 100.0%; Pred. No. 39;
Matches 15; Conservative 0; Mismatches 0; Indels 0; Gaps 0;

```

```

Qy      39  GAACCCCTGCTGATG 53
Db      20  GAACCCCTGCTGATG 34

```

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RESULT 14
US-09-736-457-1590
; Sequence 1590, Application US/09736457
; Patent No. 6509448
; GENERAL INFORMATION:
; APPLICANT: Wang, Tongtong
; APPLICANT: Bangur, Chaitanya S.
; APPLICANT: Lodes, Michael A.
; APPLICANT: Fanger, Gary
; APPLICANT: Vedvick, Tom
; APPLICANT: Carter, Darrick
; APPLICANT: Retter, Marc
; APPLICANT: Mannion, Jane

```

```

; APPLICANT: Fan, Liqun
; APPLICANT: Wang, Aijun
; TITLE OF INVENTION: COMPOSITIONS AND METHODS FOR THE THERAPY AND
; TITLE OF INVENTION: DIAGNOSIS OF LUNG CANCER
; FILE REFERENCE: 210121.478C15
; CURRENT APPLICATION NUMBER: US/09/736,457
; CURRENT FILING DATE: 2000-12-13
; NUMBER OF SEQ ID NOS: 1864
; SOFTWARE: FASTSEQ for Windows Version 3.0
; SEQ ID NO: 1590
; LENGTH: 434
; TYPE: DNA
; ORGANISM: Homo sapiens
; FEATURE:
; NAME/KEY: misc_feature
; LOCATION: (1)...(434)
; OTHER INFORMATION: n = A,T,C or G
US-09-736-457-1590

```

```

Query Match          23.1%; Score 15; DB 4; Length 434;
Best Local Similarity 100.0%; Pred. No. 39;
Matches 15; Conservative 0; Mismatches 0; Indels 0; Gaps 0;

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```

QY 39 GAACCCCTGCTGATG 53
DB 20 GAACCCCTGCTGATG 34

```

```

RESULT 15
US-09-614-124B-1590
; Sequence 1590: Application US/09614124B
; Patent No. 6630574
; GENERAL INFORMATION:
; APPLICANT: Wang, Tonglong
; APPLICANT: Bangur, Chaitanya S.
; APPLICANT: Lodes, Michael A.
; APPLICANT: Fanger, Gary
; APPLICANT: Vedvick, Tom
; APPLICANT: Carter, Darrick
; APPLICANT: Retter, Marc
; APPLICANT: Manion, Jane
; TITLE OF INVENTION: COMPOSITIONS AND METHODS FOR THERAPY AND
; TITLE OF INVENTION: DIAGNOSIS OF LUNG CANCER
; FILE REFERENCE: 210121.478C9
; CURRENT APPLICATION NUMBER: US/09/614,124B
; CURRENT FILING DATE: 2001-07-11
; NUMBER OF SEQ ID NOS: 1668
; SOFTWARE: FASTSEQ for Windows Version 3.0
; SEQ ID NO: 1590
; LENGTH: 434
; TYPE: DNA
; ORGANISM: Homo sapiens
; FEATURE:
; NAME/KEY: misc_feature
; LOCATION: (1)...(434)
; OTHER INFORMATION: n = A,T,C or G
US-09-614-124B-1590

```

```

Query Match          23.1%; Score 15; DB 4; Length 434;
Best Local Similarity 100.0%; Pred. No. 39;
Matches 15; Conservative 0; Mismatches 0; Indels 0; Gaps 0;

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```

QY 39 GAACCCCTGCTGATG 53
DB 20 GAACCCCTGCTGATG 34

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Search completed: December 23, 2004, 01:33:34
 Job time : 55.3529 secs

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Query Match	100.0%	Score 65;	DB 9;	Length 65;
Best Local Similarity	100.0%	Pred. No. 1.3e-27;		
Matches 65;	Conservative 0;	Mismatches 0;	Indels 0;	Gaps 0;
QY	1	GATTCATGGAATATGCCCCGCTTTCCAGCGCTTTATTCGAAACCTCGTGTGACACCC	60	
DB	1	GATTCATGGAATATGCCCCGCTTTCCAGCGCTTTATTCGAAACCTCGTGTGACACCC	60	
OY	61	CGTCC	65	
DB	61	CGTCC	65	

RESULT 2

US-09-898-616A-1
Sequence 1, Application US/09898616A
Publication No. US20030109429A1
GENERAL INFORMATION:
APPLICANT: Claragen Inc.
APPLICANT: Pilon, Aprile L
APPLICANT: Welch, Richard W
TITLE OF INVENTION: Method for the Production of Purified rhug for the Treatment of
TITLE OF INVENTION: Inflammatory and Fibrotic Conditions
FILE REFERENCE: 116142/00170
CURRENT APPLICATION NUMBER: US/09/898,616A
PRIOR FILING DATE: 2002-10-15
PRIOR APPLICATION NUMBER: US 08/864,357
PRIOR FILING DATE: 1997-05-28
NUMBER OF SEQ ID NOS: 10
SOFTWARE: PatentIn version 3.1
SEQ ID NO 1
LENGTH: 65
TYPE: DNA
ORGANISM: Artificial Sequence
FEATURE:
OTHER INFORMATION: Original amino acid sequence obtained from cDNA. Current submitte
FEATURE:
NAME/KEY: misc feature
OTHER INFORMATION: Original amino acid sequence obtained from cDNA. Current submitte
OTHER INFORMATION: d sequence maximized for expression in E. coli.
US-09-898-616A-1

Query Match

Best Local Similarity 100.0%; Score 65; DB 10; Length 65;
Best Local Similarity 100.0%; Pred. No. 1.3e-27;
Matches 65; Conservative 0; Mismatches 0; Indels 0; Gaps 0;

QY 1 GATCCATGGAATCTGCGCTTTCCAGCGTGTATCGAAACCGTGTATGACACCC 60

DB 1 GATCCATGGAATCTGCGCTTTCCAGCGTGTATCGAAACCGTGTATGACACCC 60

QY 61 CGTCC 65

DB 61 CGTCC 65

RESULT 3

US-10-187-498A-1
Sequence 1, Application US/10187498A
Publication No. US2003020795A1
GENERAL INFORMATION:
APPLICANT: Claragen Inc.
APPLICANT: Pilon, Aprile L
APPLICANT: Welch, Richard W
TITLE OF INVENTION: Method for the Production of Purified rhug for the Treatment of
TITLE OF INVENTION: Inflammatory and Fibrotic Conditions
FILE REFERENCE: 116142/00260
CURRENT APPLICATION NUMBER: US/10/187,498A
PRIOR FILING DATE: 2001-07-02
PRIOR APPLICATION NUMBER: US 08/864,357
PRIOR FILING DATE: 1997-05-28
NUMBER OF SEQ ID NOS: 10
SOFTWARE: PatentIn version 3.1
SEQ ID NO 1
LENGTH: 65
TYPE: DNA
ORGANISM: Artificial Sequence
FEATURE:
OTHER INFORMATION: Original amino acid sequence obtained from cDNA. Current submitte
OTHER INFORMATION: d sequence maximized for expression in E. coli.
NAME/KEY: misc feature
OTHER INFORMATION: Original amino acid sequence obtained from cDNA. Current submitte
OTHER INFORMATION: d sequence maximized for expression in E. coli.

US-10-187-498A-1

Query Match 100.0%; Score 65; DB 15; Length 65;
Best Local Similarity 100.0%; Pred. No. 1.3e-27;
Matches 65; Conservative 0; Mismatches 0; Indels 0; Gaps 0;

QY 1 GATCCATGGAATCTGCGCTTTCCAGCGTGTATCGAAACCGTGTATGACACCC 60

DB 1 GATCCATGGAATCTGCGCTTTCCAGCGTGTATCGAAACCGTGTATGACACCC 60

QY 61 CGTCC 65

DB 61 CGTCC 65

RESULT 4

US-10-647-371-5
Sequence 5, Application US/10647371
Publication No. US20040047857A1
GENERAL INFORMATION:
APPLICANT: Claragen, Inc. & NIH
TITLE OF INVENTION: Use of Recombinant Human Uteroglobulin in Treatment of Inflammatory
TITLE OF INVENTION: and Fibrotic Conditions
FILE REFERENCE: 116142-85
CURRENT APPLICATION NUMBER: US/10/647,371
PRIOR FILING DATE: 2003-08-25
PRIOR APPLICATION NUMBER: 09/549,926
PRIOR FILING DATE: 2000-04-14
NUMBER OF SEQ ID NOS: 12
SOFTWARE: PatentIn version 3.2
SEQ ID NO 5
LENGTH: 65
TYPE: DNA
ORGANISM: Artificial
FEATURE:
OTHER INFORMATION: Description of Artificial Sequence: primer sequence
US-10-647-371-5

Query Match 100.0%; Score 65; DB 15; Length 65;
Best Local Similarity 100.0%; Pred. No. 1.3e-27;
Matches 65; Conservative 0; Mismatches 0; Indels 0; Gaps 0;

QY 1 GATCCATGGAATCTGCGCTTTCCAGCGTGTATCGAAACCGTGTATGACACCC 60

DB 1 GATCCATGGAATCTGCGCTTTCCAGCGTGTATCGAAACCGTGTATGACACCC 60

QY 61 CGTCC 65

DB 61 CGTCC 65

RESULT 5

US-09-861-688-13/c
Sequence 13, Application US/09861688
Patent No. US2002017460A1
GENERAL INFORMATION:
APPLICANT: Claragen, Inc. & NIH
TITLE OF INVENTION: Use of Recombinant Human Uteroglobulin in Treatment of
TITLE OF INVENTION: Inflammatory and
TITLE OF INVENTION: Fibrotic Conditions
FILE REFERENCE: 116142/2
CURRENT APPLICATION NUMBER: US/09/861,688
PRIOR FILING DATE: 2001-05-21
PRIOR APPLICATION NUMBER: 08/864,357
PRIOR FILING DATE: 1997-05-28
NUMBER OF SEQ ID NOS: 22
SOFTWARE: PatentIn version 3.0
SEQ ID NO 13
LENGTH: 42
TYPE: DNA
ORGANISM: Artificial Sequence
FEATURE:
OTHER INFORMATION: primer sequence

US-09-861-688-13

Query Match
Best Local Similarity 100.0%; Pred. No. 3.1e-14;
Matches 42; Conservative 0; Mismatches 0; Indels 0; Gaps 0;

QY 1 GATCCATGGAATCTGCCCGCTTTTCAGCGCTTTATCGAA 42
DB 42 GATCCATGGAATCTGCCCGCTTTTCAGCGCTTTATCGAA 1

RESULT 6
US-10-647-371-12/c

/ Sequence 12, Application US/10647371
/ Publication No. US20040047857A1
/ GENERAL INFORMATION:
/ APPLICANT: Clargen, Inc. & NIH
/ TITLE OF INVENTION: Use of Recombinant Human Uteroglobulin in Treatment of Inflammatory
/ TITLE OF INVENTION: and Fibrotic Conditions
/ FILE REFERENCE: 116142-85
/ CURRENT APPLICATION NUMBER: US/10/647,371
/ PRIOR FILING DATE: 2003-08-25
/ PRIOR APPLICATION NUMBER: 09/549,926
/ PRIOR FILING DATE: 2000-04-14
/ NUMBER OF SEQ ID NOS: 12
/ SOFTWARE: PatentIn version 3.2
/ SEQ ID NO 12
/ LENGTH: 42
/ TYPE: DNA
/ ORGANISM: Artificial
/ FEATURE:
/ OTHER INFORMATION: Description of Artificial Sequence: primer sequence
US-10-647-371-12

Query Match
Best Local Similarity 100.0%; Pred. No. 3.1e-14;
Matches 42; Conservative 0; Mismatches 0; Indels 0; Gaps 0;

QY 1 GATCCATGGAATCTGCCCGCTTTTCAGCGCTTTATCGAA 42
DB 42 GATCCATGGAATCTGCCCGCTTTTCAGCGCTTTATCGAA 1

RESULT 7
US-09-861-688-12/c

/ Sequence 12, Application US/09861688
/ Patent No. US20020173460A1
/ GENERAL INFORMATION:
/ APPLICANT: Clargen, Inc. & NIH
/ TITLE OF INVENTION: Use of Recombinant Human Uteroglobulin in Treatment of
/ TITLE OF INVENTION: Inflammatory and
/ TITLE OF INVENTION: Fibrotic Conditions
/ FILE REFERENCE: 116142/2
/ CURRENT APPLICATION NUMBER: US/09/861,688
/ CURRENT FILING DATE: 2001-05-21
/ PRIOR APPLICATION NUMBER: 08/864,357
/ PRIOR FILING DATE: 1997-05-28
/ NUMBER OF SEQ ID NOS: 22
/ SOFTWARE: PatentIn version 3.0
/ SEQ ID NO 12
/ LENGTH: 60
/ TYPE: DNA
/ ORGANISM: Artificial Sequence
/ FEATURE:
/ OTHER INFORMATION: primer sequence
US-09-861-688-12

Query Match
Best Local Similarity 100.0%; Pred. No. 0.0034;
Matches 23; Conservative 0; Mismatches 0; Indels 0; Gaps 0;

QY 43 CCTGCTGATGAGACCCCGTCC 65
|||||

DB 60 CCTGCTGATGAGACCCCGTCC 38

RESULT 8
US-09-898-616A-7/c

/ Sequence 7, Application US/09898616A
/ Publication No. US20030109429A1
/ GENERAL INFORMATION:
/ APPLICANT: Clargen Inc.
/ APPLICANT: Pilon, Apple L
/ TITLE OF INVENTION: Method for the Production of Purified rhNG for the Treatment of
/ TITLE OF INVENTION: Inflammatory and Fibrotic Conditions
/ FILE REFERENCE: 116142/00170
/ CURRENT APPLICATION NUMBER: US/09/898,616A
/ CURRENT FILING DATE: 2002-10-15
/ PRIOR APPLICATION NUMBER: US 08/864,357
/ PRIOR FILING DATE: 1997-05-28
/ NUMBER OF SEQ ID NOS: 10
/ SOFTWARE: PatentIn version 3.1
/ SEQ ID NO 7
/ LENGTH: 60
/ TYPE: DNA
/ ORGANISM: Artificial Sequence
/ FEATURE:
/ OTHER INFORMATION: Original amino acid sequence obtained from cDNA. Current submitte
/ OTHER INFORMATION: d sequence maximized for expression in E. coli.
/ NAME/KEY: misc feature
/ OTHER INFORMATION: Original amino acid sequence obtained from cDNA. Current submitte
US-09-898-616A-7

Query Match
Best Local Similarity 100.0%; Pred. No. 0.0034;
Matches 23; Conservative 0; Mismatches 0; Indels 0; Gaps 0;

QY 43 CCTGCTGATGAGACCCCGTCC 65
DB 60 CCTGCTGATGAGACCCCGTCC 38

RESULT 9
US-09-898-616A-8/c

/ Sequence 8, Application US/09898616A
/ Publication No. US20030109429A1
/ GENERAL INFORMATION:
/ APPLICANT: Clargen Inc.
/ APPLICANT: Pilon, Apple L
/ TITLE OF INVENTION: Method for the Production of Purified rhNG for the Treatment of
/ TITLE OF INVENTION: Inflammatory and Fibrotic Conditions
/ FILE REFERENCE: 116142/00170
/ CURRENT APPLICATION NUMBER: US/09/898,616A
/ CURRENT FILING DATE: 2002-10-15
/ PRIOR APPLICATION NUMBER: US 08/864,357
/ PRIOR FILING DATE: 1997-05-28
/ NUMBER OF SEQ ID NOS: 10
/ SOFTWARE: PatentIn version 3.1
/ SEQ ID NO 8
/ LENGTH: 60
/ TYPE: DNA
/ ORGANISM: Artificial Sequence
/ FEATURE:
/ OTHER INFORMATION: Original amino acid sequence obtained from cDNA. Current submitte
/ OTHER INFORMATION: d sequence maximized for expression in E. coli.
/ NAME/KEY: misc feature
/ OTHER INFORMATION: Original amino acid sequence obtained from cDNA. Current submitte
US-09-898-616A-8

Query Match
Best Local Similarity 100.0%; Pred. No. 0.0034;
Matches 23; Conservative 0; Mismatches 0; Indels 0; Gaps 0;

Best Local Similarity 100.0%; Pred. No. 0.0034;
Matches 23; Conservative 0; Mismatches 0; Indels 0; Gaps 0;

QY 43 CCTGCTGATGAGACCCCGTCC 65
DB 60 CCTGCTGATGAGACCCCGTCC 38

RESULT 10

US-10-187-498A-7/C

Sequence 7, Application US/10187498A
Publication No. US2003020795A1

GENERAL INFORMATION:

APPLICANT: Clargen Inc.

APPLICANT: Pilon, Aprile L

APPLICANT: Welch, Richard W

TITLE OF INVENTION: Method for the Production of Purified rhug for the Treatment of

TITLE OF INVENTION: Inflammatory and Fibrotic Conditions

FILE REFERENCE: 116142/00260

CURRENT FILING DATE: 2001-07-02

PRIOR APPLICATION NUMBER: US 08/864,357

PRIOR FILING DATE: 1997-05-28

NUMBER OF SEQ ID NOS: 10

SOFTWARE: PatentIn version 3.1

SEQ ID NO 7

LENGTH: 60

TYPE: DNA

ORGANISM: Artificial Sequence

FEATURE: OTHER INFORMATION: Original amino acid sequence obtained from cDNA. Current submitte

FEATURE: OTHER INFORMATION: d sequence maximized for expression in E. coli.

NAME/KEY: misc feature

OTHER INFORMATION: Original amino acid sequence obtained from cDNA. Current submitte

OTHER INFORMATION: d sequence maximized for expression in E. coli.

US-10-187-498A-7

Query Match 35.4%; Score 23; DB 15; Length 60;
Best Local Similarity 100.0%; Pred. No. 0.0034;
Matches 23; Conservative 0; Mismatches 0; Indels 0; Gaps 0;

QY 43 CCTGCTGATGAGACCCCGTCC 65
DB 60 CCTGCTGATGAGACCCCGTCC 38

RESULT 11

US-10-187-498A-8/C

Sequence 8, Application US/10187498A
Publication No. US2003020795A1

GENERAL INFORMATION:

APPLICANT: Clargen Inc.

APPLICANT: Pilon, Aprile L

APPLICANT: Welch, Richard W

TITLE OF INVENTION: Method for the Production of Purified rhug for the Treatment of

TITLE OF INVENTION: Inflammatory and Fibrotic Conditions

FILE REFERENCE: 116142/00260

CURRENT FILING DATE: 2001-07-02

PRIOR APPLICATION NUMBER: US 08/864,357

PRIOR FILING DATE: 1997-05-28

NUMBER OF SEQ ID NOS: 10

SOFTWARE: PatentIn version 3.1

SEQ ID NO 8

LENGTH: 60

TYPE: DNA

ORGANISM: Artificial Sequence

FEATURE: OTHER INFORMATION: Original amino acid sequence obtained from cDNA. Current submitte

FEATURE: OTHER INFORMATION: d sequence maximized for expression in E. coli.

NAME/KEY: misc feature

OTHER INFORMATION: Original amino acid sequence obtained from cDNA. Current submitte
OTHER INFORMATION: d sequence maximized for expression in E. coli.
US-10-187-498A-8

Query Match 35.4%; Score 23; DB 15; Length 60;
Best Local Similarity 100.0%; Pred. No. 0.0034;
Matches 23; Conservative 0; Mismatches 0; Indels 0; Gaps 0;

QY 43 CCTGCTGATGAGACCCCGTCC 65
DB 60 CCTGCTGATGAGACCCCGTCC 38

RESULT 12

US-10-647-371-11/C

Sequence 11, Application US/10647371
Publication No. US20040047857A1

GENERAL INFORMATION:

APPLICANT: Clargen, Inc. & NIH

TITLE OF INVENTION: Use of Recombinant Human Uteroglobulin in Treatment of Inflammatory

TITLE OF INVENTION: and Fibrotic Conditions

FILE REFERENCE: 116142-85

CURRENT FILING DATE: 2003-08-25

PRIOR APPLICATION NUMBER: 09/549,926

PRIOR FILING DATE: 2000-04-14

NUMBER OF SEQ ID NOS: 12

SOFTWARE: PatentIn version 3.2

SEQ ID NO 11

LENGTH: 60

TYPE: DNA

ORGANISM: Artificial

FEATURE: OTHER INFORMATION: Description of Artificial Sequence: primer sequence

US-10-647-371-11

Query Match 35.4%; Score 23; DB 16; Length 60;
Best Local Similarity 100.0%; Pred. No. 0.0034;
Matches 23; Conservative 0; Mismatches 0; Indels 0; Gaps 0;

QY 43 CCTGCTGATGAGACCCCGTCC 65
DB 60 CCTGCTGATGAGACCCCGTCC 38

RESULT 13

US-10-129-422-10

Sequence 10, Application US/10129422
Publication No. US2003016251A1

GENERAL INFORMATION:

APPLICANT: KIM, Sun Young et al.; Wyomed Ltd.

TITLE OF INVENTION: High efficiency retroviral vector which contains genetically engi

TITLE OF INVENTION: non-coding sequence harboring splice acceptor

FILE REFERENCE: PCA10935/VML/PCT

CURRENT FILING DATE: 2002-10-16

NUMBER OF SEQ ID NOS: 15

SOFTWARE: KOPATIN 1.5

SEQ ID NO 10

LENGTH: 36

TYPE: DNA

ORGANISM: Artificial Sequence

FEATURE: OTHER INFORMATION: primer IRAP5'

US-10-129-422-10

Query Match 26.2%; Score 17; DB 15; Length 36;
Best Local Similarity 100.0%; Pred. No. 10;
Matches 17; Conservative 0; Mismatches 0; Indels 0; Gaps 0;

QY 1 GATCCATGGAATCTGC 17
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Db 2 GATCCATGGAATCTGC 18

RESULT 14

US-10-425-115-112555
 ; Sequence 112555, Application US/10425115
 ; Publication No. US20040214272A1
 ; GENERAL INFORMATION:
 ; APPLICANT: La Rosa, Thomas J.
 ; APPLICANT: Kovalic, David K.
 ; APPLICANT: Zhou, Yihua
 ; APPLICANT: Cao, Yongwei
 ; TITLE OF INVENTION: Nucleic Acid Molecules and Other Molecules Associated With
 ; FILE REFERENCE: 38-21(53222)B
 ; CURRENT APPLICATION NUMBER: US/10/425,115
 ; NUMBER OF SEQ ID NOS: 369326
 ; SEQ ID NO 112555
 ; LENGTH: 534
 ; TYPE: DNA
 ; ORGANISM: Zea mays
 ; FEATURE:
 ; OTHER INFORMATION: Clone ID: MRT4577_34141C.1
 US-10-425-115-112555

Query Match 26.2%; Score 17; DB 18; Length 534;
 Best Local Similarity 100.0%; Pred. No. 9.7;
 Matches 17; Conservative 0; Mismatches 0; Indels 0; Gaps 0;

QY 12 ATCTGCCCGCTTTCCA 28
 DB 454 ATCTGCCCGCTTTCCA 470

RESULT 15

US-10-424-599-87144
 ; Sequence 87144, Application US/10424599
 ; Publication No. US20040031072A1
 ; GENERAL INFORMATION:
 ; APPLICANT: La Rosa, Thomas J.
 ; APPLICANT: Kovalic, David K.
 ; APPLICANT: Zhou, Yihua
 ; APPLICANT: Cao, Yongwei
 ; TITLE OF INVENTION: Soy Nucleic Acid Molecules and Other Molecules Associated With
 ; FILE REFERENCE: 38-21(53223)B
 ; CURRENT APPLICATION NUMBER: US/10/424,599
 ; CURRENT FILING DATE: 2003-04-28
 ; NUMBER OF SEQ ID NOS: 285684
 ; SEQ ID NO 87144
 ; LENGTH: 647
 ; TYPE: DNA
 ; ORGANISM: Glycine max
 ; FEATURE:
 ; OTHER INFORMATION: Clone ID: PAT_MRT3847_49701C.1
 US-10-424-599-87144

Query Match 26.2%; Score 17; DB 16; Length 647;
 Best Local Similarity 100.0%; Pred. No. 9.7;
 Matches 17; Conservative 0; Mismatches 0; Indels 0; Gaps 0;

QY 33 GTTATCGAAACCTGCT 49
 DB 413 GTTATCGAAACCTGCT 429

Search completed: December 23, 2004, 05:19:24
 Job time: 920.177 secs

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GenCore version 5.1.6
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OM nucleic - nucleic search, using sw model

Run on: December 22, 2004, 19:17:58 ; Search time 848.688 Seconds
(without alignments)
3343.258 Million cell updates/sec

Title: US-09-898-616A-2

Perfect score: 60

Sequence: 1 agctacgaagcagctatgta.....acatgcgtgaagcaggtgct 60

Scoring table: OLIGO_NUC
Gapop 60.0 , Gapext 60.0

Searched: 4526729 seqs, 23644849745 residues

Word size : 0

Total number of hits satisfying chosen parameters: 3053458

Minimum DB seq length: 0

Maximum DB seq length: 2000000000

Post-processing: Listing first 45 summaries

Database :

GenEmbl1:*
1: gb_ba:*
2: gb_hlg:*
3: gb_in:*
4: gb_om:*
5: gb_ov:*
6: gb_pat:*
7: gb_ph:*
8: gb_pl:*
9: gb_pr:*
10: gb_ro:*
11: gb_sts:*
12: gb_sy:*
13: gb_un:*
14: gb_vi:*

Pred. No. is the number of results predicted by chance to have a score greater than or equal to the score of the result being printed, and is derived by analysis of the total score distribution.

SUMMARIES

Result No.	Score	Query Match	Length	DB ID	Description
1	60	100.0	60	6	ARI60916
2	37	61.7	60	6	ARI60916 Sequence
3	19	31.7	89959	8	AP004967
4	19	31.7	200849	2	AC026895
5	18	30.0	59	6	ARI60920
6	18	30.0	201	11	BV203590
7	18	30.0	201	11	BV203591
8	18	30.0	201	11	BV203592
9	18	30.0	432	6	AX397069
10	18	30.0	1428	6	BD094070
11	18	30.0	1428	9	HUMSF2P33
12	18	30.0	1618	5	BC042354
13	18	30.0	1717	9	HUMASF
14	18	30.0	1860	5	BC067552
15	18	30.0	2002	5	BC055511
16	18	30.0	2055	5	BC075558
17	18	30.0	2059	5	BC033785
18	18	30.0	2369	5	BC046679
19	18	30.0	2583	9	AB000463

20	18	30.0	2708	9	BC010264	BC010264 Homo sapi
21	18	30.0	2731	5	BC076945	BC076945 Xenopus t
22	18	30.0	2785	6	BD189398	BD189398 Tumor ant
23	18	30.0	2765	9	AB062124	AB062124 Homo sapi
24	18	30.0	2900	5	BC066682	BC066682 Dantio rer
25	18	30.0	22970	9	HS1247F6	268279 Human DNA s
26	18	30.0	35240	2	AC040987	AC040987 Homo sapi
27	18	30.0	69023	2	AC087445	AC087445 Homo sapi
28	18	30.0	110000	2	BX272305_0	BX272305 Homo sapi
29	18	30.0	119974	9	AC018763	AC018763 Dantio rer
30	18	30.0	135299	9	AC091857	AC091857 Homo sapi
31	18	30.0	142801	2	AC079206	AC079206 Homo sapi
32	18	30.0	149969	9	HSJ323A24	AL121750 Human DNA
33	18	30.0	154462	2	AC036136	AC036136 Homo sapi
34	18	30.0	160111	2	AP000780	AP000780 Homo sapi
35	18	30.0	160415	2	AC140961	AC140961 Papio anu
36	18	30.0	164404	2	AC027149	AC027149 Homo sapi
37	18	30.0	165726	2	AC143663	AC143663 Macaca mu
38	18	30.0	168651	9	AC009474	AC009474 Homo sapi
39	18	30.0	169403	9	AL445929	AL445929 Human DNA
40	18	30.0	176310	2	BX897686	BX897686 Dantio rer
41	18	30.0	176879	5	AL929224	AL929224 Zebrafish
42	18	30.0	177314	9	AC022209	AC022209 Homo sapi
43	18	30.0	181275	2	BX511308	BX511308 Dantio rer
44	18	30.0	181494	2	AC015813	AC015813 Homo sapi
45	18	30.0	196476	2	AC055890	AC055890 Homo sapi

ALIGNMENTS

RESULT 1
LOCUS ARI60916
DEFINITION Sequence 7 from patent US 6255281.
ACCESSION ARI60916
VERSION ARI60916.1 GI:16225981
KEYWORDS
SOURCE
ORGANISM
REFERENCE
AUTHORS
TITLE
JOURNAL
FEATURES
source

Unknown.
Unclassified.
1 (bases 1 to 60)

Pilon A.L., Mukherjee, A.B. and Zhang, Z.
Use of recombinant human uteroglobin in treatment of inflammatory
and fibrotic conditions
Patent: US 6255281-A 7 03-JUN-2001;
Location/Qualifiers

1..60
/organism="unknown"
/mol_type="unassigned DNA"

ORIGIN

Query Match 100.0%; Score 60; DB 6; Length 60;
Best Local Similarity 100.0%; Pred. No. 1.4e-22;
Matches 60; Conservative 0; Mismatches 0; Indels 0; Gaps 0;

QY 1 AGCTACGAAGCAGCTATGGAATCTCTCCGACACGACATCGTGAAGCAGGTGCT 60
DB 1 AGCTACGAAGCAGCTATGGAATCTCTCCGACACGACATCGTGAAGCAGGTGCT 60

RESULT 2

ARI60921/c
LOCUS ARI60921
DEFINITION Sequence 12 from patent US 6255281.
ACCESSION ARI60921
VERSION ARI60921.1 GI:16225996
KEYWORDS
SOURCE
ORGANISM
REFERENCE
AUTHORS

Unknown.
Unclassified.
1 (bases 1 to 60)
Pilon, A.L., Mukherjee, A.B. and Zhang, Z.

TITLE Use of recombinant human uteroglobin in treatment of inflammatory and fibrotic conditions
 JOURNAL Patent: US 6255281-A 12 03-JUL-2001;
 FEATURES Location/Qualifiers
 source 1..60
 /organism="unknown"
 /mol_type="unassigned DNA"

ORIGIN
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 Best Local Similarity 100.0%; Pred. No. 1,1e-09;
 Matches 37; Conservative 0; Mismatches 0; Indels 0; Gaps 0;

QY 1 AGCTGAGAGAGCTATGAACTGTTCTCCGAGAC 37
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 37 AGCTGAGAGAGCTATGAACTGTTCTCTCCGAGAC 1

Db

RESULT 3
 AP004967/c 89959 bp DNA linear PLN 22-JUL-2003
 LOCUS Lotus corniculatus var. japonicus genomic DNA, chromosome 1,
 DEFINITION clone:UJT27L02, TM0144, complete sequence.
 ACCESSION AP004967
 VERSION AP004967.1 GI:21907985
 KEYWORDS HTG.
 SOURCE Lotus corniculatus var. japonicus (Lotus japonicus)
 ORGANISM Lotus corniculatus var. japonicus
 Eukaryota; Viridiplantae; Streptophyta; Embryophyta; Tracheophyta;
 Spermatophyta; Magnoliophyta; eudicotyledons; core eudicots;
 rosids; eurosids I; Fabales; Fabaceae; Papilionoideae; Loteeae;
 Lotus.

REFERENCE
 AUTHORS 1
 TITLE Kaneko, T., Nakamura, Y., Asamizu, E., Kato, T., Sato, S. and Tabata, S.
 Structural Analysis of a Lotus japonicus Genome. I. Sequence
 Features and Mapping of Sixty-six TAC clones which cover the 6.7 Mb
 Regions of the Genome
 Regions of the Genome
 Unpublished
 2 (bases 1 to 89959)
 JOURNAL
 REFERENCE
 AUTHORS Nakamura, Y.
 TITLE Direct Submission
 JOURNAL Submitted (26-MAR-2002) Yasukazu Nakamura, Kazusa DNA Research
 Institute, Department of Plant Gene Research, 1532-3, Yata,
 Kisarazu, Chiba 292-0812, Japan (E-mail: ynakazusa.or.jp,
 URL: http://www.kazusa.or.jp; Tel: 81-438-52-3935,
 Fax: 81-438-52-3934)

FEATURES
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 Location/Qualifiers
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 Best Local Similarity 100.0%; Pred. No. 6;
 Matches 19; Conservative 0; Mismatches 0; Indels 0; Gaps 0;

QY 8 AAGCAGCTATGAACTGTT 26
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 Db 87202 AAGCAGCTATGAACTGTT 87184

RESULT 4
 AC026895 200849 bp DNA linear HTG 20-APR-2000
 LOCUS Homo sapiens chromosome 2 clone RP11-396J2 map 2, WORKING DRAFT
 DEFINITION
 SEQUENCE 22 unordered pieces.
 AC026895

VERSION
 KEYWORDS
 SOURCE
 ORGANISM

REFERENCE
 AUTHORS
 TITLE
 JOURNAL
 REFERENCE
 AUTHORS

AC026895.2 GI:7596820
 HTG; HTGS PHASE1; HTGS_DRAFT.
 Homo sapiens (human)
 Homo sapiens
 Eukaryota; Metazoa; Chordata; Craniata; Vertebrata; Euteleostomi;
 Mammalia; Eutheria; Primates; Catarrhini; Homnidae; Homo.
 1 (bases 1 to 200849)
 Birren, B., Linton, L., Nusbaum, C. and Lander, E.
 Homo sapiens chromosome 2, clone RP11-396J2
 Unpublished
 2 (bases 1 to 200849)
 Birren, B., Linton, L., Nusbaum, C., Lander, E., Abraham, H., Allen, N.,
 Anderson, S., Baldwin, J., Barna, N., Bastien, V., Bedalov, F.,
 Boguski, M., Brown, A., Brown, A., Burkett, G.,
 Campion, A., Castle, A., Chao, Y., Collins, S.,
 Collymore, A., Cooke, P., DeRubeis, K., Dewar, K., Diaz, J., S.,
 Dodge, S., Domingo, M., Doyle, M., Ferreira, P., Fitzhugh, W., Gage, D.,
 Galagan, J., Gardy, S., Glendon, S., Goyette, M., Graham, L.,
 Grand, P., Grant, G., Hagan, B., Heaford, A., Horton, L.,
 Howland, J., Iley, I., Johnson, R., Jones, C., Kan, L., Karas, A.,
 Klein, J., Lacroque, K., Lamazares, R., Landers, T., Lech, J.,
 Levine, R., Liu, C., Liu, G., Locke, K., MacDonald, P., Margis, N.,
 McCarthy, M., McEwan, P., McGurk, A., McKernan, K., McPherson, R.,
 Meldrum, J., Meneses, L., Mihov, T., Miranda, C., Miska, V., Morrow, J.,
 Murphy, T., Naylor, J., Norman, C., O'Connor, T., O'Donnell, P.,
 O'Neill, D., Oliver, T., Oliver, J., Peterson, K., Pierre, N.,
 Pisani, C., Pollara, V., Raymond, C., Riley, R., Rogov, P., Rothman, D.,
 Roy, A., Santos, R., Schauer, S., Severy, P., Spencer, B.,
 Stange-Thomann, N., Stojanovic, N., Subramanian, A., Talamas, J.,
 Tassile, S., Theodore, J., Tirrell, A., Travers, M., Triggillo, J.,
 Vassiliev, H., Viel, R., Vo, A., Wilson, B., Wu, X., Wyman, D., Ye, W., J.,
 Young, G., Zainoun, J., Zimmer, A. and Zody, M.
 Direct Submission
 Submitted (25-MAR-2000) Whitehead Institute/MIT Center for Genome
 Research, 320 Charles Street, Cambridge, MA 02141, USA
 On Apr 19, 2000 this sequence version replaced gi:7328755.
 All repeats were identified using RepeatMasker:
 Smit, A.F.A. & Green, P. (1996-1997)
 http://ftp.genome.washington.edu/JM/RepeatMasker.html

Genome Center
 Center: Whitehead Institute/ MIT Center for Genome Research
 Center code: WIBR
 Web site: http://www-seg.wi.mit.edu
 Contact: sequence_submissions@genome.wi.mit.edu
 Project Information
 Center project name: L6461
 Center clone name: 396 J 2
 Summary Statistics
 Sequencing vector: MJ3, M7815, 100% of reads
 Chemistry: Dye-terminator Big Dye, 100% of reads
 Assembly program: Phrap, version 0.960731
 Consensus quality: 188437 bases at least Q40
 Consensus quality: 194805 bases at least Q30
 Consensus quality: 197132 bases at least Q20
 Insert size: 210000; agarose-ff
 Insert size: 198749; sum-of-coverage
 Quality coverage: 4.6 in Q20 bases; agarose-ff
 Quality coverage: 4.8 in Q20 bases; sum-of-coverage

NOTE: This is a 'working draft' sequence. It currently
 consists of 22 contigs. The true order of the pieces
 is not known and their order in this sequence record as
 arbitrary. Gaps between the contigs are represented as
 runs of N, but the exact sizes of the gaps are unknown.
 This record will be updated with the finished sequence
 as soon as it is available and the accession number will
 be preserved.

1 1067: contig of 1067 bp in length
 * 1 1068 1167: gap of 100 bp
 * 1 1168 2886: contig of 1689 bp in length
 * 1 2887 2956: gap of 100 bp
 * 1 2957 5983: contig of 3027 bp in length
 * 1 5984 6083: gap of 100 bp

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* 6084 7783: contig of 1700 bp in length
* 7784 7883: gap of 100 bp
* 7884 10615: contig of 2732 bp in length
* 10616 10715: gap of 100 bp
* 10716 12692: contig of 1977 bp in length
* 12693 12792: gap of 100 bp
* 12793 14959: contig of 2167 bp in length
* 14960 15059: gap of 100 bp
* 15060 19764: contig of 4705 bp in length
* 19765 19864: gap of 100 bp
* 19865 22667: contig of 2802 bp in length
* 22667 27132: contig of 4366 bp in length
* 27133 31434: contig of 4202 bp in length
* 31435 31534: gap of 100 bp
* 31535 39440: contig of 7906 bp in length
* 39441 46513: gap of 100 bp
* 46514 46613: contig of 673 bp in length
* 46614 55509: contig of 8896 bp in length
* 55510 66257: gap of 10647 bp in length
* 66257 7020: contig of 10664 bp in length
* 7021 77120: gap of 100 bp
* 77121 88050: contig of 10930 bp in length
* 88051 88150: gap of 100 bp
* 88151 98020: contig of 9870 bp in length
* 98021 98120: gap of 100 bp
* 98121 108280: contig of 10160 bp in length
* 108281 108380: gap of 100 bp
* 108381 126228: contig of 17848 bp in length
* 126229 126328: gap of 100 bp
* 126329 165333: contig of 38905 bp in length
* 165334 165334: gap of 100 bp
* 165334 200849: contig of 35516 bp in length.

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FEATURES
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/clone_id="RPC1-11 Human Male BAC"
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Matches 19; Conservative 0; Mismatches 0; Indels 0; Gaps 0;
QY 9 AGCAGCTATGAGACTGTTTC 27
Db 114115 AGCAGCTATGAGACTGTTTC 114133

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RESULT 5
LOCUS AR160920/c 59 bp DNA linear PAT 17-OCT-2001
DEFINITION Sequence 11 from patent US 6255281.
ACCESSION AR160920
VERSION AR160920.1 GI:16225993
KEYWORDS
SOURCE Unknown.
ORGANISM Unknown.
REFERENCE 1 (bases 1 to 59)
AUTHORS Pilon,A.L., Mukherjee,A.B. and Zhang,Z.
TITLES Use of recombinant human uteroglobin in treatment of inflammatory
and fibrotic conditions
JOURNAL Patent: US 6255281-A 11 03-JUL-2001;
FEATURES
LOCATION/Qualifiers
1..59
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ORIGIN
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Best Local Similarity 100.0%; Pred. No. 4.7;
Matches 18; Conservative 0; Mismatches 0; Indels 0; Gaps 0;
QY 38 AGACATCGTGAAGCAG 55
Db 59 AGACATCGTGAAGCAG 42

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RESULT 6
LOCUS BV203590 201 bp DNA linear STS 10-JUN-2004
DEFINITION sqm212590 Human DNA (sequenc) Homo sapiens STS genomic, sequence
tagged site.
ACCESSION BV203590
VERSION BV203590.1 GI:48173010
KEYWORDS STS.
SOURCE Homo sapiens (human)
ORGANISM Homo sapiens
Eukaryota; Metazoa; Chordata; Craniata; Vertebrata; Euteleostomi;

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REFERENCE
1 (bases 1 to 201)
Mammalia; Eutheria; Primates; Catarrhini; Homidae; Homo.

AUTHORS
Nelson, R.M., Marnellos, G., Kammerer, S., Hoyal, C.R., Shi, M.M.,
Cantor, C.R. and Braun, A.
TITLE
Large-Scale Validation of Single Nucleotide Polymorphisms in Gene
Regions
JOURNAL
Genome Res. (2004) In press

COMMENT
Contact: Andreas Braun
Pharmaceuticals division
Sequenom, Inc.
3595 John Hopkins Court, San Diego, CA 92121, USA
Tel: 18582029018
Fax: 18582029020
Email: abraun@sequenom.com

Primer A: No primer sequence submitted
Primer B: No primer sequence submitted
STS size: 201.

FEATURES
source
1..201
Location/Qualifiers

ORIGIN
STS
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/clone_lib="Human DNA (Sequenom)"
<1..>201

Query Match
Best Local Similarity 100.0%; Pred. No. 42;
Matches 18; Conservative 0; Mismatches 0; Indels 0; Gaps 0;

QY
41 ACATGCGTGAAGCAGGTG 58
130 ACATGCGTGAAGCAGGTG 147

RESULT 7
BV203591 201 bp DNA linear STS 10-JUN-2004
DEFINITION
sqm212591 Human DNA (Sequenom) Homo sapiens STS genomic, sequence
tagged site.

ACCESSION
BV203591
VERSION
BV203591.1 GI:48173011
KEYWORDS
STS.

SOURCE
Homo sapiens (human)
ORGANISM
Homo sapiens
Eukaryota; Metazoa; Chordata; Craniata; Vertebrata; Euteleostomi;
Mammalia; Eutheria; Primates; Catarrhini; Homidae; Homo.
REFERENCE
1 (bases 1 to 201)
Nelson, R.M., Marnellos, G., Kammerer, S., Hoyal, C.R., Shi, M.M.,
Cantor, C.R. and Braun, A.
TITLE
Large-Scale Validation of Single Nucleotide Polymorphisms in Gene
Regions
JOURNAL
Genome Res. (2004) In press

COMMENT
Contact: Andreas Braun
Pharmaceuticals division
Sequenom, Inc.
3595 John Hopkins Court, San Diego, CA 92121, USA
Tel: 18582029018
Fax: 18582029020
Email: abraun@sequenom.com

Primer A: No primer sequence submitted
Primer B: No primer sequence submitted
STS size: 201.

FEATURES
source
1..201
Location/Qualifiers

ORIGIN
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/clone_lib="Human DNA (Sequenom)"
<1..>201

Query Match
Best Local Similarity 100.0%; Pred. No. 42;
Matches 18; Conservative 0; Mismatches 0; Indels 0; Gaps 0;

QY
41 ACATGCGTGAAGCAGGTG 58
121 ACATGCGTGAAGCAGGTG 138

RESULT 8
BV203592 201 bp DNA linear STS 10-JUN-2004
DEFINITION
sqm212593 Human DNA (Sequenom) Homo sapiens STS genomic, sequence
tagged site.

ACCESSION
BV203592
VERSION
BV203592.1 GI:48173012
KEYWORDS
STS.

SOURCE
Homo sapiens (human)
ORGANISM
Homo sapiens
Eukaryota; Metazoa; Chordata; Craniata; Vertebrata; Euteleostomi;
Mammalia; Eutheria; Primates; Catarrhini; Homidae; Homo.
REFERENCE
1 (bases 1 to 201)
Nelson, R.M., Marnellos, G., Kammerer, S., Hoyal, C.R., Shi, M.M.,
Cantor, C.R. and Braun, A.
TITLE
Large-Scale Validation of Single Nucleotide Polymorphisms in Gene
Regions
JOURNAL
Genome Res. (2004) In press

COMMENT
Contact: Andreas Braun
Pharmaceuticals division
Sequenom, Inc.
3595 John Hopkins Court, San Diego, CA 92121, USA
Tel: 18582029018
Fax: 18582029020
Email: abraun@sequenom.com

Primer A: No primer sequence submitted
Primer B: No primer sequence submitted
STS size: 201.

FEATURES
source
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Location/Qualifiers

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/clone_lib="Human DNA (Sequenom)"
<1..>201

Query Match
Best Local Similarity 100.0%; Pred. No. 42;
Matches 18; Conservative 0; Mismatches 0; Indels 0; Gaps 0;

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112 ACATGCGTGAAGCAGGTG 129

RESULT 9
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LOCUS
Sequence 1284 from Patent WO0212328.
DEFINITION
AX397069
ACCESSION
AX397069.1 GI:21067816
VERSION
AX397069.1
KEYWORDS
STS.

SOURCE
Homo sapiens (human)
ORGANISM
Homo sapiens
Eukaryota; Metazoa; Chordata; Craniata; Vertebrata; Euteleostomi;
Mammalia; Eutheria; Primates; Catarrhini; Homidae; Homo.

REFERENCE
1
King, G.E., Meagher, M.J., Xu, J. and Secrist, H.
Compositions and methods for the therapy and diagnosis of colon
cancer
Patent: WO 0212328-A 1284 14-FEB-2002;

Amphibia; Batrachia; Anura; Mesobatrachia; Pipidae; Pipidae;
Xenopodinae; Xenopus; Xenopus.
1 (bases 1 to 1618)
Klein, S.L., Strausberg, R.L., Wagner, L., Pontius, J., Clifton, S.W.,
and Richardson, P.
Genetic and genomic tools for Xenopus research: The NIH Xenopus
initiative
Dev. Dyn. 225 (4), 384-391 (2002)

JOURNAL
PUBMED
REFERENCE
AUTHORS
2 (bases 1 to 1618)
Strausberg, R.L., Feingold, E.A., Grouse, L.H., Derge, J.G.,
Klausner, R.D., Collins, P.S., Wagner, L., Shenmen, C.M., Schuler, G.D.,
Holtzman, S.F., Zeeberg, B., Buetow, K.H., Schaefer, C.F., Bhat, N.K.,
Hopkins, R.F., Jordan, H., Moore, T., Max, S.I., Wang, J., Hsieh, F.,
Diatchenko, L., Mansina, K., Farmer, A.A., Rubin, G.M., Hong, L.,
Scapellato, M., Soares, M.B., Bonaldo, M.F., Casavant, T.L.,
Scheetz, T.E., Brownstein, M.J., Usdin, T.B., Toshiyuki, S.,
Carninci, P., Prange, C., Raha, S.S., Loquellano, N.A., Peters, G.J.,
Abramson, R.D., Mullany, S.J., Bosak, S.A., McEwan, P.J.,
McKernan, K.J., Malek, J.A., Gunaratne, P.H., Richards, S.,
Morley, K.C., Hale, S., Garcia, A.M., Gay, L.J., Hulik, S.W.,
Villalón, D.K., Muzny, D.M., Sodergren, E.J., Lu, X., Gibbs, R.A.,
Raeey, J., Helton, E., Kettelman, M., Madan, A., Rodriguez, S.,
Sanchez, A., Whiting, M., Madan, A., Young, A.C., Shevchenko, Y.,
Boutard, G.G., Blakesley, R.W., Touchman, J.W., Green, E.D.,
Dickson, M.C., Rodriguez, A.C., Grimwood, J., Schmutz, J., Myers, R.M.,
Butterfield, Y.S., Krzywinski, M.I., Skalska, U., Smal, D.E.,
Schneringer, A., Schein, J.E., Jones, S.J., and Marra, M.A.
Generation and initial analysis of more than 15,000 full-length
human and mouse cDNA sequences
Proc. Natl. Acad. Sci. U.S.A. 99 (26), 16899-16903 (2002)

JOURNAL
PUBMED
REFERENCE
AUTHORS
JOURNAL
TITLE
3 (bases 1 to 1618)
Klein, S. and Strausberg, R.
Direct Submission
Submitted (02-JAN-2003) National Institutes of Health, Xenopus Gene
Collection (XGC), National Institute of Child Health and Human
Development, 6100 Executive Boulevard, Room 4B01, Rockville, MD
20892-7510, USA
NIH-XGC Project
Contact: XGC help desk
Email: xgcs-help@nih.gov
Tissue Procurement: Dr. Igor David
CNA Library Preparation: Life Technologies, Inc.
CNA Library Arrayed by: The I.M.A.G.E. Consortium (LLNL)
DNA Sequencing by: Institute for Systems Biology
http://www.systemsbio.org
Contact: amadasystemsbio.org
Anup Madan, Jessica Fahey, Erin Helton, Mark Kettelman, Anuradha
Madan, Stephanie Rodrigues, Amy Sanchez and Michelle Whiting

REMARK
COMMENT
NIH-XGC Project
Contact: XGC help desk
Email: xgcs-help@nih.gov
Tissue Procurement: Dr. Igor David
CNA Library Preparation: Life Technologies, Inc.
CNA Library Arrayed by: The I.M.A.G.E. Consortium (LLNL)
DNA Sequencing by: Institute for Systems Biology
http://www.systemsbio.org
Contact: amadasystemsbio.org
Anup Madan, Jessica Fahey, Erin Helton, Mark Kettelman, Anuradha
Madan, Stephanie Rodrigues, Amy Sanchez and Michelle Whiting

FEATURES
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of an RNA binding protein. RBMs are found in a variety of
RNA binding proteins, including various hnRNP proteins,
proteins implicated in regulation of alternative splicing,
and protein components of snRNPs. The motif also appears
in a few single stranded DNA binding proteins. The RBM
structure consists of four strands and two helices
arranged in an alpha/beta sandwich, with a third helix
present during RNA binding in some cases. The C-terminal
beta strand (4th strand) and final helix are hard to align
and have been omitted in the SEED alignment. The LA
proteins have a N terminus rim which is aligned in the
seed. There is a second region towards the C terminus that
has some features of a rim but does not appear to have the
important structural core of a rim. The LA proteins are
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Matches 18; Conservative 0; Mismatches 0; Indels 0; Gaps 0;
QY 41 ACATGCGTGAAGCAGGTG 58
Db 525 ACATGCGTGAAGCAGGTG 542

RESULT 13
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LOCUS 1717 bp mRNA linear PRI 27-APR-1993
DEFINITION Human alternative splicing factor mRNA, complete cds.
ACCESSION M72709.1
VERSION M72709.1 GI:179073
KEYWORDS alternative splicing factor.
SOURCE Homo sapiens (human)
ORGANISM Homo sapiens
Eukaryota; Metazoa; Chordata; Craniata; Vertebrata; Euteleostomi;
Mammalia; Eutheria; Primates; Catarrhini; Homnidae; Homo.
1 (bases 1 to 1717)
Ge, H., Zuo, P. and Manley, J.L.
Primary structure of the human splicing factor ASF reveals
similarities with Drosophila regulators
Cell 66 (2), 373-382 (1991)

REFERENCE
AUTHORS
TITLE
JOURNAL
MEDLINE
PUBMED
COMMENT
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mRNA


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Matches 18; Conservative 0; Mismatches 0; Indels 0; Gaps 0;

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RESULT 14
LOCUS      BC056752                1860 bp    mRNA    linear    VRT 25-MAY-2004
DEFINITION Dantio rerio cDNA clone MGC:65898 IMAGE:6801570, complete cds.
ACCESSION  BC056752
VERSION     BC056752.1  GI:34785173
KEYWORDS    MGC.
SOURCE      Dantio rerio (zebrafish)
ORGANISM    Dantio rerio
            Metazoa; Chordata; Craniata; Vertebrata; Euteleostomi;
            Actinopterygii; Neopterygii; Teleostei; Ostariophysi;
            Cypriniformes; Cyprinidae; Dantio.
            1 (bases 1 to 1860)
            Strausberg R.L., Feingold E.A., Grouse L.H., Derge J.G.,
            Klausner R.D., Collins F.S., Wagner L., Shenmen C.M., Schuler G.D.,
            Altschul S.F., Zeeberg B., Buetow K.H., Scheffer C.F., Bhat N.K.,
            Hopkins R.F., Jordan H., Moore T., Max S.I., Wang J., Hsieh F.,
            Datchenko L., Marusina K., Farmer A.A., Rubin G.M., Hong L.,
            Stajich M., Soares M.B., Bonaldi M.F., Casavant T.L.,
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            Carninci P., Prange C., Raha S.S., Loquellano N.A., Peters G.J.,

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TITLE
JOURNAL
PUBMED
REFERENCE
AUTHORS
JOURNAL
TITLE
2 (bases 1 to 1860)
Strausberg, R.
Direct Submission
Submitted (25-AUG-2003) National Institutes of Health, Mammalian
Gene Collection (MGC), Cancer Genomics Office, National Cancer
Institute, 31 Center Drive, Room 11A03, Bethesda, MD 20892-2590,
USA
NIH-MGC Project URL: http://mgc.nci.nih.gov
Contact: MGC help desk
Email: cgabs-r@mail.nih.gov
Tissue Procurement: Dr. Chi-Bin Chien
CDNA Library Preparation: Invitrogen Corp
CDNA Library Arrayed by: The I.M.A.G.E. Consortium (LNL)
DNA Sequencing by: National Institutes of Health Intramural
Sequencing Center (NISC),
Gaithersburg, Maryland;
Web site: http://www.nisc.nih.gov/
Contact: nisc.mgc@nih.gov
Aklter N., Ayle K., Beckstrom-Sternberg S.M., Benjamin B.,
Blakesley R.W., Bouffard G.G., Bren K., Brinkley C., Brooks S.,
Dietrich N.L., Granite S., Guan X., Gupta V., Haghighi P.,
Hansen N., Ho S.-L., Karlins E., Kwong P., Latic P., Legaspi R.,
Maduro Q.L., Mastello C., Maskeri B., Mastrian S.D., McCloskey J.C.,
McDowell J., Pearson R., Stancijop S., Thomas P.D., Touchman J.W.,
Tsurgenc C., Vogt J.L., Walker M.A., Weherby K.D., Wiggins L.,
Young A., Zhang L.-H. and Green E.D.
Clone distribution: MGC clone distribution information can be found
through the I.M.A.G.E. Consortium/LNLW at: http://image.llnl.gov
Series: IRXK Plate: 121 Row: P Column: 12
This clone was selected for full length sequencing because it
passed the following selection criteria: Hexamer frequency ORF
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Location/Qualifiers
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Best Local Similarity 100.0%; Pred. No. 33;
Matches 18; Conservative 0; Mismatches 0; Indels 0; Gaps 0;

CY      41 ACATGCGTGAAGCAGGTG 58

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Db 611 ACATGCGTGAACGAGGTG 628

RESULT 15 BC055511

LOCUS BC055511 2002 bp mRNA linear VRT 30-JUN-2004
DEFINITION Danio rerio zgc:66146, mRNA (cDNA clone MGC:66146 IMAGE:5410436), complete cds.

ACCESSION BC055511
VERSION BC055511.1 GI:33416586
KEYWORDS MGC.
SOURCE Danio rerio (zebrafish)
ORGANISM Danio rerio

REFERENCE 1 (bases 1 to 2002)
Strausberg, R.L., Feingold, E.A., Grouse, L.H., Derge, J.G., Klausner, R.D., Collins, F.S., Wagner, L., Shenmen, C.M., Schuler, G.D., Altschul, S.F., Zeeberg, B., Buetow, K.H., Schaefer, C.F., Bhat, N.K., Hopkins, R.F., Jordan, H., Moore, T., Max, S.I., Wang, J., Hsieh, F., Diatchenko, L., Mansina, K., Farmer, A.A., Rubin, G.M., Hong, L., Stupp, S., Soares, M.B., Sontag, M.F., Casavant, T.L., Scheetz, T.E., Brownstein, M.U., Usdin, T.B., Tashy, S., Carninci, P., Prange, C., Raha, S.S., Loguercio, N.A., Peters, G.J., Abramson, R.D., Mullahy, S.J., Bosak, S.A., McEwan, P.J., McKernan, K.J., Malek, J.A., Gunaratne, P.H., Richards, S., Worley, K.C., Hale, S., Garcia, A.M., Gay, L.J., Hult, S.W., Villalón, D.K., Muzny, D.M., Sodergren, E.J., Lu, X., Gibbs, R.A., Fahey, J., Helton, E., Kettman, M., Madan, A., Rodriguez, S., Sanchez, A., Whiting, M., Madan, A., Young, A.C., Shevchenko, Y., Bouffard, G.G., Blakesley, R.W., Touchman, J.W., Green, E.D., Dickson, M.C., Rodriguez, A.C., Grimwood, J., Schmutz, J., Myers, R.M., Butlerfield, Y.S., Krzywinski, M.I., Skalska, U., Small, D.E., Scherch, A., Schein, J.E., Jones, S.J., and Marra, M.A.
Generation and initial analysis of more than 15,000 full-length human and mouse cDNA sequences
Proc Natl. Acad. Sci. U.S.A. 99 (26), 16699-16903 (2002)

JOURNAL 12477932
PUBMED 2 (bases 1 to 2002)
AUTHORS Strausberg, R.
TITLE Direct Submission
JOURNAL Submitted (01-AUG-2003) National Institutes of Health, Mammalian Gene Collection (MGC), Cancer Genomics Office, National Cancer Institute, 31 Center Drive, Room 11A03, Bethesda, MD 20892-2590, USA

REMARK NIE-MGC Project URL: <http://mgc.nci.nih.gov>
COMMENT Contact: MGC help desk
Email: cgaps-remail.nih.gov
Tissue Procurement: Dr. Sumio Sugano
CDNA Library Preparation: Dr. Sumio Sugano
CDNA Library Arrayed by: The I.M.A.G.E. Consortium (LITL)
DNA Sequencing by: Sequencing Group at the Stanford Human Genome Center, Stanford University School of Medicine, Stanford, CA 94305
Web site: <http://www.shgc.stanford.edu>
Contact: (Dickson, Mark) mcd@paxil.stanford.edu
Dickson, M., Schmutz, J., Grimwood, J., Rodriguez, A., and Myers, R. M.

FEATURES
source
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/organism="Danio rerio"
/mol_type="mRNA"
/db_xref="taxon:7955"
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DKNRGGRPRFVFRERPDADAVYARQYVDGRLVFRPRSRGMRGRGFRGG
GGGGGGGGGGGAFGRGTFPSRSRIRIVSGLPSSGMDLKHMRADGVCA
DVFRDGTVEVFRKEDYIVAKLDNFKPSHGETAYIRKVDGSPSPYGRSRR
SRSRSRSRSRNNRSYSPPRSRGGSP"

ORIGIN
Query Match 30.0%; Score 18; DB 5; Length 2002;
Best Local Similarity 100.0%; Pred. No. 33;
Matches 18; Conservative 0; Mismatches 0; Indels 0; Gaps 0;

CY 41 ACATGCGTGAACGAGGTG 58
DB 528 ACATGCGTGAACGAGGTG 545

Search completed: December 22, 2004, 23:36:31
Job time : 654.668 secs

GenCore version 5.1.6
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OM nucleic - nucleic search, using sw model

Run on: December 22, 2004, 19:14:28 ; Search time 211.765 Seconds
(without alignments)
1457.336 Million cell updates/sec

Title: US-09-898-616a-2

Perfect score: 60
Sequence: 1 agctacgaagcagcgtcgtga.....acatgcgtgaagcagcgtcgtc 60

Scoring table: OLIGO_NUC
Gapop 60.0, Gapext 60.0

Searched: 4134886 seqs, 2624710521 residues

Word size: 0

Total number of hits satisfying chosen parameters: 8269772

Minimum DB seq length: 0

Maximum DB seq length: 2000000000

Post-processing: Listing first 45 summaries

Database:

N_Geneseq_23sep04:*
1: geneseqn1980s:*
2: geneseqn1890s:*
3: geneseqn2000s:*
4: geneseqn2001s:*
5: geneseqn2001bs:*
6: geneseqn2002as:*
7: geneseqn2002bs:*
8: geneseqn2003as:*
9: geneseqn2003bs:*
10: geneseqn2003cs:*
11: geneseqn2003ds:*
12: geneseqn2004s:*

Pred. No. is the number of results predicted by chance to have a score greater than or equal to the score of the result being printed, and is derived by analysis of the total score distribution.

SUMMARIES

Result No.	Score	Query Match	Length	ID	Description
1	60	100.0	60	9	ABZ58371
2	60	100.0	60	12	ADL27627
3	37	61.7	60	9	ABZ58376
4	37	61.7	60	9	ABZ58377
5	37	61.7	60	12	ADL27633
6	37	61.7	60	12	ADL27632
7	37	61.7	60	12	ADL27632
8	37	61.7	60	12	ADL27632
9	37	61.7	60	12	ADL27632
10	37	61.7	60	12	ADL27632
11	37	61.7	60	12	ADL27632
12	37	61.7	60	12	ADL27632
13	37	61.7	60	12	ADL27632
14	37	61.7	60	12	ADL27632
15	37	61.7	60	12	ADL27632
16	37	61.7	60	12	ADL27632
17	37	61.7	60	12	ADL27632
18	37	61.7	60	12	ADL27632
19	37	61.7	60	12	ADL27632
20	37	61.7	60	12	ADL27632
21	37	61.7	60	12	ADL27632

ALIGNMENTS

22	17	28.3	1978	6	ABN59992	ABN59992 Novel hum
23	16	26.7	65	6	ABN56135	ABN56135 Mouse spl
24	16	26.7	300	2	AAZ14356	AAZ14356 Human gen
25	16	26.7	351	12	ADL85790	ADL85790 DNA up-re
26	16	26.7	351	12	ADL85789	ADL85789 DNA up-re
27	16	26.7	373	8	ABX50810	ABX50810 Bovine ES
28	16	26.7	624	9	ADA30641	ADA30641 DNA encod
29	16	26.7	732	2	AAZ17762	AAZ17762 Human gen
30	16	26.7	802	2	AAZ16289	AAZ16289 Human gen
31	16	26.7	830	11	AD131178	AD131178 Human CDN
32	16	26.7	846	3	AAZ49282	AAZ49282 Arabidops
33	16	26.7	869	6	ABK13152	ABK13152 Transcript
34	16	26.7	1019	4	AAZ59766	AAZ59766 Pseudonib
35	16	26.7	1019	8	ACF64695	ACF64695 Pseudonib
36	16	26.7	1820	10	ADJ18824	ADJ18824 Human dis
37	16	26.7	1919	11	ADM03518	ADM03518 Human CDN
38	16	26.7	1959	11	ABD09379	ABD09379 Pseudonib
39	16	26.7	2010	11	ABD09565	ABD09565 Pseudonib
40	16	26.7	2402	4	AAK81667	AAK81667 Human imm
41	16	26.7	3602	8	ACF12861	ACF12861 Human cer
42	16	26.7	3803	2	AAZ76396	AAZ76396 Human sec
43	16	26.7	3803	10	ADC38821	ADC38821 Human CDN
44	16	26.7	5333	10	ADD47032	ADD47032 Human gen
45	16	26.7	5333	10	ADD47860	ADD47860 Human gen

RESULT 1
ID ABZ58371 standard, DNA, 60 BP.
AC ABZ58371;
DT 28-APR-2003 (first entry)
XX Human uteroglobin synthet gene oligonucleotide 2.
DE Human, uteroglobin, respiratory distress; antiinflammatory; antifibrotic;
KW antiinflammatory; antiaesthetic; nephrotropic; antineuritic;
XX antitartaric; ss.
XX Homo sapiens.
OS Synthetic.
XX MO2003003979-A2.
XX 16-JAN-2003.
XX 02-JUL-2002; 2002WO-US020836.
XX 02-JUL-2001; 2001US-00898616.
XX (CLAR-) CLARAGEN INC.
XX Pilon AL, Welch RE;
XX WPI; 2003-221527/21.
XX Bacterial expression system for producing recombinant human uteroglobin
PT for treating inflammatory and fibrotic conditions, comprises a synthetic
PT gene which codes for human uteroglobin.
XX Claim 1; Page 33; 127pp; English.
XX The present sequence is that of oligonucleotide 2, which was used in the
XX construction of a synthetic gene for the production of human uteroglobin
XX (hug) in bacteria. Oligonucleotides 1-4 (see ABZ58370-73) were used to
XX assemble the coding strand and oligonucleotides 5-8 (see ABZ58374-77) the
XX complementary strand. The gene was assembled by annealing and ligation of
XX the oligonucleotides. Because mature native hug has glutamic acid at its
XX N-terminus, an initiator methionine was added to the N-terminus, and

CC codon usage was optimised for expression in bacteria. In an example from
CC the invention, the synthetic gene was cloned into plasmid pCG12 (see
CC AB258378) and recombinant hUG (see ABP72259) was produced in *Escherichia*
CC coli strain CG12. The invention relates generally to the production of
CC recombinant hUG by bacterial expression, protein purification practices, the
CC up production according to current good manufacturing practices, the
CC recombinant hUG is useful for the treatment of inflammatory and fibrotic
CC conditions, such as neonatal respiratory distress syndrome and
CC bronchopulmonary dysplasia. It may also be used to treat conditions
CC associated with elevated phospholipase A2 levels such as pancreatitis,
CC acute renal failure, rheumatoid arthritis and asthma

SO Sequence 60 BP; 15 A; 15 C; 18 G; 12 T; 0 U; 0 Other;

Query Match 100.0%; Score 60; DB 9; Length 60;
Best Local Similarity 100.0%; Pred. No. 5.2e-23;
Matches 60; Conservative 0; Mismatches 0; Indels 0; Gaps 0;

DB 1 AGCTACGAGCAGCTATGGAAGTCTCTCCGACGACGACATGCGTGAAGCAGGTGCT 60
1 AGCTACGAGCAGCTATGGAAGTCTCTCTCCGACGACGACATGCGTGAAGCAGGTGCT 60

RESULT 2
ADL27627 standard; DNA; 60 BP.

AC ADL27627;
XX 20-MAY-2004 (first entry)
DT
XX Recombinant human uteroglobin, rhUG, coding oligonucleotide #2.
DE
XX Human; ss; recombinant human uteroglobin; rhUG;
KM bacterial expression system; rhUG master cell bank;
KW rhUG research seed bank; anti-inflammatory; secretory phospholipase A 2;
KW fibronectin; respiratory distress; inflammation; fibrotic disease.

XX Homo sapiens.
OS Synthetic.
XX US2003207795-A1.
PN 06-NOV-2003.
PD 02-JUL-2002; 2002US-00187498.
PF 28-MAY-1997; 97US-00864357.
PR 02-JUL-2001; 2001US-00898616.
XX (PILON) PILON A L.
PA (WELCH) WELCH R W.
PI Pilon AL, Welch RW;
XX WPI; 2004-051527/05.
DR Bacterial expression system for production of recombinant human
XX uteroglobin comprising synthetic gene or human cDNA sequence which codes
PT for human uteroglobin.
XX Claim 1; SEQ ID NO 2; 64pp; English.

XX The invention relates to a bacterial expression system for the production
CC of recombinant human uteroglobin (rhUG), comprising a synthetic gene or
CC human cDNA sequence which codes for human UG, constructed from the
CC oligonucleotides appearing as ADL27626-ADL27629, and which further
CC comprises Met-Ala-Ala at the N-terminus of the sequence. Also included
CC are producing an rhUG master cell bank (comprising inoculating a suitable
CC incubating broth with an aliquot portion of a rhUG research seed bank to
CC form a bacterial culture, incubating the bacterial culture, adding a
CC cryopreservative to the bacterial culture to form a cryopreserved
CC solution, transferring a portion of the cryopreserved solution to a

CC cryovial and storing the cryovial at a temperature below -60 degrees C),
CC expressing rhUG (comprising providing a production seed cell bank culture
CC comprising an expression vector capable of expressing rhUG, inoculating a
CC broth medium with the production seed cell bank culture to form an
CC inoculum, incubating the bacterial culture formed in step (b),
CC inoculating a large scale fermenter with the inoculum formed from the
CC step (c) to form a fermentation culture, adding an induction agent to
CC culture within the large scale fermenter, adding an induction agent to
CC the fermentation culture to induce the expression of rhUG and harvesting
CC the above fermentation culture), purifying rhUG, determining the potency
CC of rhUG in a sample, measuring in vitro anti-inflammatory effect arising
CC from inhibition or blocking of secretory phospholipase A 2 enzymes by
CC rhUG, measuring in vitro binding of rhUG to fibronectin, determining the
CC purity of rhUG, and a pharmaceutical composition comprising a purified
CC rhUG and a carrier or diluent. The bacterial expression system is useful
CC for producing a rhUG research seed bank or a pharmaceutical grade rhUG
CC drug substance. rhUG is safe to administer to a patient in respiratory
CC distress. The rhUG is useful for treating inflammation and fibrotic
CC diseases. The present sequence is a coding strand oligonucleotide used to
CC construct the synthetic rhUG gene.

SO Sequence 60 BP; 15 A; 15 C; 18 G; 12 T; 0 U; 0 Other;

Query Match 100.0%; Score 60; DB 12; Length 60;
Best Local Similarity 100.0%; Pred. No. 5.2e-23;
Matches 60; Conservative 0; Mismatches 0; Indels 0; Gaps 0;

DB 1 AGCTACGAGCAGCTATGGAAGTCTCTCCGACGACGACATGCGTGAAGCAGGTGCT 60
1 AGCTACGAGCAGCTATGGAAGTCTCTCTCCGACGACGACATGCGTGAAGCAGGTGCT 60

RESULT 3
ABZ58376/c
ID ABZ58376 standard; DNA; 60 BP.

XX ABZ58376;
AC 28-APR-2003 (first entry)
DT
XX Human uteroglobin synthetic gene oligonucleotide 7.
DE
XX Human; uteroglobin; respiratory distress; anti-inflammatory; antifibrotic;
KM anti-inflammatory; antiaesthetic; nephrotoxic; anti-neumatic;
KW antiarthritic; ss.
XX Homo sapiens.
OS Synthetic.
XX WO2003003979-A2.
PN 16-JAN-2003.
PD 02-JUL-2002; 2002WMO-US020836.
PF 02-JUL-2001; 2001US-00898616.
XX (CLAR-) CLARAGEN INC.
PA Pilon AL, Welch RE;
XX WPI; 2003-221527/21.
DR Bacterial expression system for producing recombinant human uteroglobin
XX for treating inflammatory and fibrotic conditions, comprises a synthetic
XX gene which codes for human uteroglobin.
XX Example 1; Page 33; 127pp; English.

XX The present sequence is that of oligonucleotide 7, which was used in the
CC construction of a synthetic gene for the production of human uteroglobin
CC (rhUG) in bacteria. Oligonucleotides 1-4 (see ABZ58370-73) were used to
CC assemble the coding strand and oligonucleotides 5-8 (see ABZ58374-77) the

complementary strand. The gene was assembled by annealing and ligation of the oligonucleotides. Because mature native hUG has glutamic acid at its N-terminus, an initiator methionine was added to the N-terminus, and codon usage was optimised for expression in bacteria. In an example from the invention, the synthetic gene was cloned into plasmid pCG12 (see AB258378) and recombinant hUG (see ABP72259) was produced in *Escherichia coli* strain CG12. The invention relates generally to the production of recombinant hUG by bacterial expression, protein purification and scaled-up production according to current good manufacturing practices. The recombinant hUG is useful for the treatment of inflammatory and fibrotic conditions, such as neonatal respiratory distress syndrome and bronchopulmonary dysplasia. It may also be used to treat conditions associated with elevated phospholipase A2 levels such as pancreatitis, acute renal failure, rheumatoid arthritis and asthma.

Sequence 60 BP; 12 A; 14 C; 22 G; 12 T; 0 U; 0 Other;

Query Match 61.7%; Score 37; DB 9; Length 60;
Best Local Similarity 100.0%; Pred. No. 2.1e-10;
Matches 37; Conservative 0; Mismatches 0; Indels 0; Gaps 0;

1 AGCTACGAGCAGCTATGCACTGTTCTCTCCGAGCC 37
37 AGCTACGAGCAGCTATGCACTGTTCTCTCCGAGCC 1

RESULT 4
AB258377/c
ID AB258377 standard; DNA; 60 BP.
XX
AC AB258377;
XX
DT 28-APR-2003 (first entry)
XX
DE Human uteroglobin synthetic gene oligonucleotide 8.
XX
XX Human; uteroglobin; respiratory distress; antiinflammatory; antifibrotic;
KM antiinflammatory; antiasthmatic; nephroprotective; antirheumatic;
KM antiarthritic; ss.
XX
OS Homo sapiens.
XX Synthetic.
XX
PN WC003003979-A2.
XX
PD 16-JAN-2003.
XX
PF 02-JUL-2002; 2002WC-US020836.
XX
PR 02-JUL-2001; 2001US-00898616.
XX
PA (CLAR-) CLARAGEN INC.
XX
PI Pilon AL, Welch RE;
XX
DR WPI; 2003-221527/21.
XX
PT Bacterial expression system for producing recombinant human uteroglobin
PT for treating inflammatory and fibrotic conditions, comprises a synthetic
PT gene which codes for human uteroglobin.
XX
PS Example 1; Page 33; 127pp; English.
XX
CC The present sequence is that of oligonucleotide 8, which was used in the
CC construction of a synthetic gene for the production of human uteroglobin
CC (hUG) in bacteria. Oligonucleotides 1-4 (see AB258370-73) were used to
CC assemble the coding strand and oligonucleotides 5-8 (see AB258374-77) the
CC complementary strand. The gene was assembled by annealing and ligation of
CC the oligonucleotides. Because mature native hUG has glutamic acid at its
CC N-terminus, an initiator methionine was added to the N-terminus, and
CC codon usage was optimised for expression in bacteria. In an example from
CC the invention, the synthetic gene was cloned into plasmid pCG12 (see
CC AB258378) and recombinant hUG (see ABP72259) was produced in *Escherichia*

coli strain CG12. The invention relates generally to the production of recombinant hUG by bacterial expression, protein purification and scaled-up production according to current good manufacturing practices. The recombinant hUG is useful for the treatment of inflammatory and fibrotic conditions, such as neonatal respiratory distress syndrome and bronchopulmonary dysplasia. It may also be used to treat conditions associated with elevated phospholipase A2 levels such as pancreatitis, acute renal failure, rheumatoid arthritis and asthma.

Sequence 60 BP; 12 A; 14 C; 22 G; 12 T; 0 U; 0 Other;

Query Match 61.7%; Score 37; DB 9; Length 60;
Best Local Similarity 100.0%; Pred. No. 2.1e-10;
Matches 37; Conservative 0; Mismatches 0; Indels 0; Gaps 0;

1 AGCTACGAGCAGCTATGCACTGTTCTCTCCGAGCC 37
37 AGCTACGAGCAGCTATGCACTGTTCTCTCCGAGCC 1

RESULT 5
ADL27633/c
ID ADL27633 standard; DNA; 60 BP.
XX
AC ADL27633;
XX
DT 20-MAY-2004 (first entry)
XX
DE Recombinant human uteroglobin, rhUG, non-coding oligonucleotide #4.
XX
XX Human; ss; recombinant human uteroglobin; rhUG;
KM bacterial expression system; rhUG master cell bank;
KM rhUG research seed bank; anti-inflammatory; secretory phospholipase A 2;
KM fibronectin; respiratory distress; inflammation; fibrotic disease.
XX
XX Homo sapiens.
XX Synthetic.
XX
PN US2003207795-A1.
XX
PD 06-NOV-2003.
XX
PF 02-JUL-2002; 2002US-00187498.
XX
PR 28-MAY-1997; 97US-00864357.
XX
PR 02-JUL-2001; 2001US-00898616.
XX
PA (PILO) PILON A L.
XX (WELC) WELCH R W.
XX
PI Pilon AL, Welch RW;
XX
DR WPI; 2004-051527/05.
XX
PT Bacterial expression system for production of recombinant human
PT uteroglobin comprising synthetic gene or human cDNA sequence which codes
PT for human uteroglobin.
XX
PS Example 1; SEQ ID NO 8; 64pp; English.
XX
CC The invention relates to a bacterial expression system for the production
CC of recombinant human uteroglobin (rhUG), comprising a synthetic gene or
CC human cDNA sequence which codes for human UG, constructed from the
CC oligonucleotides appearing as ADL27626-ADL27629, and which further
CC comprises Met-Ala-Ala at the N-terminus of the sequence. Also included
CC are producing an rhUG master cell bank (comprising inoculating a suitable
CC incubating broth with an aliquot portion of a rhUG research seed bank to
CC form a bacterial culture, incubating the bacterial culture, adding a
CC cryoprotective to the bacterial culture to form a cryopreserved
CC solution, transferring a portion of the cryopreserved solution to a
CC cryovial and storing the cryovial at a temperature below -60 degrees C),
CC expressing rhUG (comprising providing a production seed cell bank culture
CC comprising an expression vector capable of expressing rhUG, inoculating a

broth medium with the production seed cell bank culture to form an inoculum, incubating the bacterial culture formed in step (b) in a large scale fermenter with the inoculum formed from the step (c) to form a fermentation culture, adding an induction agent to the fermentation culture to induce the expression of rhug and harvesting the above fermentation culture, purifying rhug, determining the potency of rhug in a sample, measuring in vitro anti-inflammatory effect arising from inhibition or blocking of secretory phospholipase A 2 enzymes by rhug, measuring in vitro binding of rhug to fibronectin, determining the purity of rhug, and a pharmaceutical composition comprising a purified rhug and a carrier or diluent. The bacterial expression system is useful for producing a rhug research seed bank or a pharmaceutical grade rhug drug substance. rhug is safe to administer to a patient in respiratory distress. The rhug is useful for treating inflammation and fibrotic diseases. The present sequence is a non-coding strand oligonucleotide used to construct the synthetic rhug gene.

Sequence 60 BP; 12 A; 14 C; 22 G; 12 T; 0 U; 0 Other;

Query Match 61.7%; Score 37; DB 12; Length 60; Best Local Similarity 100.0%; Pred. No. 2.1e-10; Matches 37; Conservative 0; Mismatches 0; Indels 0; Gaps 0;

1 AGCTACGAGCAGCTATGGAAGTCTCTCCGACC 37
37 AGCTACGAGCAGCTATGGAAGTCTCTCCGACC 1

RESULT 6
ADL27632/c
ID ADL27632 standard; DNA; 60 BP.

ADL27632;
20-MAY-2004 (first entry)

Recombinant human uteroglobin, rhug, non-coding oligonucleotide #3.

Human; ss; recombinant human uteroglobin; rhug;
bacterial expression system; rhug master cell bank;
rhug research seed bank; anti-inflammatory; secretory phospholipase A 2;
fibronectin; respiratory distress; inflammation; fibrotic disease.

Homo sapiens.
Synthetic.

US2003207795-A1.

06-NOV-2003.

02-JUL-2002; 2002US-00187498.

28-MAY-1997; 97US-00864357.

02-JUL-2001; 2001US-00898616.

(PILLO/) PILON A L.
(WELC/) WELCH R W.

Pilon AL, Welch RW;
WPI; 2004-031527/05.

Bacterial expression system for production of recombinant human uteroglobin comprising synthetic gene or human cDNA sequence which codes for human uteroglobin.

Example 1; SEQ ID NO 7; 64bp; English.

The invention relates to a bacterial expression system for the production of recombinant human uteroglobin (rhug), comprising a synthetic gene or human cDNA sequence which codes for human rhug, constructed from the oligonucleotides appearing as ADL27626-ADL27629, and which further

comprises Met-Ala-Ala at the N-terminus of the sequence. Also included are producing an rhug master cell bank (comprising inoculating a suitable incubating broth with an aliquot portion of a rhug research seed bank to form a bacterial culture, incubating the bacterial culture, adding a cryoprotective to the bacterial culture to form a cryopreserved solution, transferring a portion of the cryopreserved solution to a cryovial and storing the cryovial at a temperature below -60 degrees C), expressing rhug (comprising providing a production seed cell bank culture comprising an expression vector capable of expressing rhug, inoculating a broth medium with the production seed cell bank culture to form an inoculum, incubating the bacterial culture formed in step (b), inoculating a large scale fermenter with the inoculum formed from the step (c) to form a fermentation culture, adding an induction agent to the culture within the large scale fermenter, adding an induction agent to the fermentation culture to induce the expression of rhug and harvesting the above fermentation culture), purifying rhug, determining the potency of rhug in a sample, measuring in vitro anti-inflammatory effect arising from inhibition or blocking of secretory phospholipase A 2 enzymes by rhug, measuring in vitro binding of rhug to fibronectin, determining the purity of rhug, and a pharmaceutical composition comprising a purified rhug and a carrier or diluent. The bacterial expression system is useful for producing a rhug research seed bank or a pharmaceutical grade rhug drug substance. rhug is safe to administer to a patient in respiratory distress. The rhug is useful for treating inflammation and fibrotic diseases. The present sequence is a non-coding strand oligonucleotide used to construct the synthetic rhug gene.

Sequence 60 BP; 12 A; 14 C; 22 G; 12 T; 0 U; 0 Other;

Query Match 61.7%; Score 37; DB 12; Length 60; Best Local Similarity 100.0%; Pred. No. 2.1e-10; Matches 37; Conservative 0; Mismatches 0; Indels 0; Gaps 0;

1 AGCTACGAGCAGCTATGGAAGTCTCTCCGACC 37
37 AGCTACGAGCAGCTATGGAAGTCTCTCCGACC 1

RESULT 7
ABZ58375/c
ID ABZ58375 standard; DNA; 59 BP.

ABZ58375;
28-APR-2003 (first entry)

Human uteroglobin synthetic gene oligonucleotide 6.

Human; uteroglobin; respiratory distress; anti-inflammatory; antifibrotic;
anti-inflammatory; antiaesthetic; nephroretropic; antineumatic;
antiarthritic; ss.

Homo sapiens.
Synthetic.

WO2003003979-A2.

16-JAN-2003.

02-JUL-2002; 2002WO-US020836.

02-JUL-2001; 2001US-00898616.

(CLAR-) CLARAGEN INC.
Pilon AL, Welch RE;
WPI; 2003-221527/21.

Bacterial expression system for producing recombinant human uteroglobin for treating inflammatory and fibrotic conditions, comprising a synthetic gene which codes for human uteroglobin.

PS Example 1; Page 33; 127pp; English.

XX The present sequence is that of oligonucleotide 6, which was used in the

CC construction of a synthetic gene for the production of human uteroglobin

CC (hUG) in bacteria. Oligonucleotides 1-4 (see AB258370-73) were used to

CC assemble the coding strand and oligonucleotides 5-8 (see AB258374-77) the

CC complementary strand. The gene was assembled by annealing and ligation of

CC the oligonucleotides. Because mature native hUG has glutamic acid at its

CC N-terminus, an initiator methionine was added to the N-terminus, and

CC codon usage was optimized for expression in bacteria. In an example from

CC the invention, the synthetic gene was cloned into plasmid pCG12 (see

CC AB258378) and recombinant hUG (see ABP72259) was produced in *Escherichia*

CC coli strain CG12. The invention relates generally to the production of

CC recombinant hUG by bacterial expression, protein purification and scaled-

CC up production according to current good manufacturing practices. The

CC recombinant hUG is useful for the treatment of inflammatory and fibrotic

CC conditions, such as neonatal respiratory distress syndrome and

CC bronchopulmonary dysplasia. It may also be used to treat conditions

CC associated with elevated phospholipase A2 levels such as pancreatitis,

CC acute renal failure, rheumatoid arthritis and asthma

XX

SQ Sequence 59 BP; 9 A; 17 C; 15 G; 18 T; 0 U; 0 Other;

Query Match 30.0%; Score 18; DB 9; Length 59;

Best Local Similarity 100.0%; Pred. No. 5.5;

Matches 18; Conservative 0; Mismatches 0; Indels 0; Gaps 0;

QY 38 AGGACATGCGTGAAGCAG 55

DB 59 AGGACATGCGTGAAGCAG 42

RESULT 8

ADL27631/c

ID ADL27631 standard; DNA; 59 BP.

XX

AC ADL27631;

XX

DT 20-MAY-2004 (first entry)

DE Recombinant human uteroglobin, rhUG, non-coding oligonucleotide #2.

XX

XX Human; ss; recombinant human uteroglobin; rhUG;

XX bacterial expression system; rhUG master cell bank;

XX rhUG research seed bank; anti-inflammatory; secretory phospholipase A 2;

XX fibronectin; respiratory distress; inflammation; fibrotic disease.

XX

OS Homo sapiens.

OS Synthetic.

XX

PN US2003207795-A1.

XX

PD 06-NOV-2003.

XX

PF 02-JUL-2002; 2002US-00187498.

XX

PR 28-MAY-1997; 97US-00864357.

PR 02-JUL-2001; 2001US-00898616.

XX

PA (PILC/) PILON A L.

PA (WELC/) WELCH R W.

XX

PI Pilon AL, Welch RW;

XX

DR WPI; 2004-051527/05.

XX

PT Bacterial expression system for production of recombinant human

PT uteroglobin comprising synthetic gene or human cDNA sequence which codes

PT for human uteroglobin.

XX

PS Example 1; SEQ ID NO 6; 64pp; English.

XX

CC The invention relates to a bacterial expression system for the production

CC of recombinant human uteroglobin (rhUG), comprising a synthetic gene or

CC human cDNA sequence which codes for human UG, constructed from the

CC oligonucleotides appearing as ADL27626-ADL27629, and which further

CC comprises Met-Ala-Ala at the N-terminus of the sequence. Also included

CC are producing an rhUG master cell bank (comprising inoculating a suitable

CC incubating broth with an aliquot portion of a rhUG research seed bank to

CC form a bacterial culture, incubating the bacterial culture, adding a

CC cryoprotective to the bacterial culture to form a cryopreserved

CC solution, transferring a portion of the cryopreserved solution to a

CC cryovial and storing the cryovial at a temperature below -60 degrees C),

CC expressing rhUG (comprising providing a production seed cell bank culture

CC comprising an expression vector capable of expressing rhUG, inoculating a

CC broth medium with the production seed cell bank culture to form an

CC inoculum, incubating the bacterial culture formed in step (b),

CC inoculating a large scale fermenter with the inoculum formed from the

CC step (c) to form a fermentation culture, incubating the fermentation

CC culture within the large scale fermenter, adding an induction agent to

CC the above fermentation culture) , purifying rhUG, determining the potency

CC of rhUG in a sample, measuring in vitro anti-inflammatory effect arising

CC from inhibition or blocking of secretory phospholipase A2 enzymes by

CC rhUG, measuring in vitro binding of rhUG to fibronectin, determining the

CC purity of rhUG, and a pharmaceutical composition comprising a purified

CC rhUG and a carrier or diluent. The bacterial expression system is useful

CC for producing a rhUG research seed bank or a pharmaceutical grade rhUG

CC drug substance. rhUG is safe to administer to a patient in respiratory

CC distress. The rhUG is useful for treating inflammation and fibrotic

CC diseases. The present sequence is a non-coding strand oligonucleotide

CC used to construct the synthetic rhUG gene.

XX

SQ Sequence 59 BP; 9 A; 17 C; 15 G; 18 T; 0 U; 0 Other;

Query Match 30.0%; Score 18; DB 12; Length 59;

Best Local Similarity 100.0%; Pred. No. 5.5;

Matches 18; Conservative 0; Mismatches 0; Indels 0; Gaps 0;

QY 38 AGGACATGCGTGAAGCAG 55

DB 59 AGGACATGCGTGAAGCAG 42

RESULT 9

ABK45733

ID ABK45733 standard; cDNA; 432 BP.

XX

AC ABK45733;

XX

DT 05-JUN-2002 (first entry).

DE cDNA encoding colon tumour protein, SEQ ID NO 1284.

XX

XX Human; colon tumour; vaccine; colon cancer; immunogenic; immunotherapy;

XX gene; ss.

XX

OS Homo sapiens.

OS WO20022328-A2.

XX

PN 14-FEB-2002.

XX

PD 31-JUL-2001; 2001WO-US024218.

XX

PF 03-AUG-2000; 2000US-0222283P.

PR 28-MAR-2001; 2001US-0279763P.

PR 29-JUN-2001; 2001US-0302051P.

XX

PA (CORI-) CORIXA CORP.

XX

PI King GE, Meagher MJ, Xu J, Secret H;

XX

DR WPI; 2002-241739/29.

XX

PT New colon cancer polypeptides and polynucleotides, useful as vaccines,

PT for diagnosing, preventing, and treating colon cancer, and as markers for
PT the progression of cancer.

PS Claim 1; SEQ ID NO 1284; 147bp; English.

CC The invention relates to polynucleotides encoding colon tumour proteins.
CC The polynucleotides are encoded polypeptides are useful in pharmaceutical
CC compositions, such as vaccines, for the diagnosis, prevention, and
CC treatment of colon cancer. Polynucleotide sequences may be used as
CC hybridisation probes or primers, and in the design and preparation of
CC ribozyme molecules for inhibiting expression of tumour polypeptides and
CC proteins in tumour cells. The compositions are useful for stimulating an
CC immune response against cancer, particularly for the immunotherapy of
CC colon cancer, and as markers for the progression of cancer. ABK4450
CC ABK46237 represent coding sequences of human colon tumour proteins of the
CC invention. Note: With the exception of SEQ ID No. 1 and 2, the sequence
CC data for this patent did not form part of the printed specification but
CC was supplied by the European Patent Office

CC Sequence 432 BP; 123 A; 85 C; 103 G; 121 T; 0 U; 0 Other;

Query Match 30.0%; Score 18; DB 6; Length 432;
Best Local Similarity 100.0%; Pred. No. 5.6;
Matches 18; Conservative 0; Mismatches 0; Indels 0; Gaps 0;

QY 41 ACATGCGTGAAGCAGGTG 58
DB 9 ACATGCGTGAAGCAGGTG 26

RESULT 10
ID ACH35932 standard; cDNA; 489 BP.

AC ACH35932;

DT 13-OCT-2003 (first entry)

DE Human endothelial cell cDNA #4065.

KW Human; ss; sequencing by hybridisation; SBH; expressed sequence tag; EST;
KW genome mapping; biodiversity; genetic disorder.

OS Homo sapiens.

PN US2003073623-A1.

PD 17-APR-2003.

PF 30-JUL-2001; 2001US-00918995.

PR 30-JUL-2001; 2001US-00918995.

PA (DRMA/) DEMANAC R T.

PA (LABA/) LABAT I.

PA (STAC/) STACHE-CRAIN B.

PA (DICK/) DICKSON M C.

PA (JONE/) JONES L W.

PI Dmanac RT, Labat I, Stache-Crain B, Dickson MC, Jones LW;

DR WPI; 2003-615964/58.

PS New polynucleotide sequences obtained from various cDNA libraries, useful

PT as hybridization probes, as oligomers for PCR, for chromosome and gene

PT mapping, in the recombinant production of protein, or in generating

PT antisense DNA or RNA.

PS Claim 1; SEQ ID NO 23144; 44bp; English.

CC The invention relates to an isolated polynucleotide comprising any one of

CC 38043 cDNA sequences, appearing as ACH12789-ACH50831, whose sequence was

CC determined by the technique of SBH (sequencing by hybridisation). Also

CC included is a purified polypeptide comprising a sequence corresponding to

CC a reading frame of the novel polynucleotide. The nucleic acid sequences

CC are useful in diagnostics as expressed sequence tags (EST) for

CC identifying expressed genes or for physical mapping of the human genome,

CC in forensics, in assessing biodiversity, or in identifying mutations

CC responsible for genetic disorders and other traits. The nucleotide

CC sequences are also useful as hybridisation probes, as oligomers for PCR,

CC for chromosome and gene mapping, in the recombinant production of

CC protein, or in generating antisense DNA or RNA. The purified polypeptide

CC is one of the 38043 isolated cDNA/EST sequences. Note: The sequence data

CC for this patent did not form part of the printed specification, but was

CC obtained in electronic format directly from USPTO at

CC secdata.uspto.gov/sequence.html?ocid=20030073623

CC Sequence 489 BP; 130 A; 103 C; 134 G; 119 T; 0 U; 3 Other;

Query Match 30.0%; Score 18; DB 9; Length 489;
Best Local Similarity 100.0%; Pred. No. 5.6;
Matches 18; Conservative 0; Mismatches 0; Indels 0; Gaps 0;

QY 41 ACATGCGTGAAGCAGGTG 58
DB 122 ACATGCGTGAAGCAGGTG 139

RESULT 11
ID AAH02904 standard; DNA; 1428 BP.

AC AAH02904;

DT 15-JUN-2001 (first entry)

DE Human shear stress-response coding sequence SEQ ID NO: 61.

KW Human; shear stress-response protein; vascular disease; arteriosclerosis;

KW de.

OS Homo sapiens.

PN WO200125427-A1.

PD 12-APR-2001.

PF 02-OCT-2000; 2000WO-0P006840.

PR 01-OCT-1999; 99JP-00280976.

PA (KYOW) KYOWA HAKKO KOGYO KK.

PA (NOUI/) NOJIMA H.

PI Nojima H, Yoshieue H, Odayashi M, Ota T, Kawabata A, Sakurada K;

PI Kuga T, Sekine S, Nakamura Y, Sugano S;

DR WPI; 2001-266308/27.

DR P-PDB; AAB90781.

PT DNA sequences, proteins encoded by them and antibodies against them

PT useful in diagnosis and treatment of vascular disease caused by

PT arteriosclerosis.

PS Claim 20; Page 386-388; 678bp; Japanese.

CC The present invention provides the protein and coding sequences of a

CC number of human shear stress response proteins. These are useful in the

CC diagnosis, treatment and screening of vascular diseases caused by

CC arteriosclerosis, including heart failure, post-PTCA restenosis and

CC hypertension

CC Sequence 1428 BP; 341 A; 296 C; 400 G; 391 T; 0 U; 0 Other;

Query Match 30.0%; Score 18; DB 4; Length 1428;

Best Local Similarity 100.0%; Pred. No. 5.6;
Matches 18; Conservative 0; Mismatches 0; Indels 0; Gaps 0;

QY 41 ACATGCGTGAAGCAGGTG 58
DB 543 ACATGCGTGAAGCAGGTG 560

RESULT 12

AA160055
ID AA160055 standard; cDNA; 1428 BP.

AC AA160055;

DT 27-AUG-2003 (first entry)

DE Human PC036-2 cDNA.

XX Human; differentially regulated protein; prevention; therapy; vaccine;

KM prostate cancer; gene therapy; pre-mRNA splicing factor; PC036-2;

XX chromosome 17q21.3-q22; gene; ss.

OS Homo sapiens.

XX Key Location/Qualifiers

FT CDS 125..871

FT /tag= a

FT /product= "Human PC036-2 protein"

PD MO2003040331-A2.

XX 15-MAY-2003.

XX 07-NOV-2002; 2002MO-US035563.

XX 07-NOV-2001; 2001US-0331041P.

XX 07-NOV-2001; 2001US-0331042P.

XX 18-DEC-2001; 2001US-0340251P.

XX 07-JUN-2002; 2002US-0344791P.

XX (ORIG-) ORIGENS TECHNOLOGIES INC.

XX Sun Z, Li X, Jay G, Kovacs KF, Fan W,

XX WPI; 2003-449451/42.

XX P-PSDB; AAC29561.

XX New polynucleotide for diagnosing, staging, monitoring, prognosticating,

XX preventing or treating, or determining the predisposition to, diseases or

XX conditions such as prostate cancer, and for research or forensic science.

XX Claim 29; Page 127-128; 100bp; English.

XX The present invention relates to novel differentially regulated genes and

XX polypeptides encoded by them. Sequences of the invention are useful in

XX diagnosing, staging, monitoring, prognosticating, preventing, treating or

XX determining the predisposition to diseases or conditions such as prostate

XX cancer. They may be used as molecular markers, drug targets, vaccines, in

XX gene therapy, research, clinical medicine or forensic science. The

XX present sequence is a differentially regulated prostate cDNA, PC036-2

XX which codes for a pre-mRNA splicing factor. PC036-2 gene is located on

XX chromosome 17q21.3-q22

XX Sequence 1428 BP; 341 A; 286 C; 400 G; 391 T; 0 U; 0 Other;

XX Query Match 30.0%; Score 18; DB 10; Length 1428;

XX Best Local Similarity 100.0%; Pred. No. 5.6;

XX Matches 18; Conservative 0; Mismatches 0; Indels 0; Gaps 0;

QY 41 ACATGCGTGAAGCAGGTG 58

DB 543 ACATGCGTGAAGCAGGTG 560

RESULT 13

ADH28741
ID ADH28741 standard; DNA; 1717 BP.

AC ADH28741;

DT 11-MAR-2004 (first entry)

DE Human chronic myelogenous leukaemia (CML) gene marker #9.

XX ds; chronic phase chronic myelogenous leukaemia; CP-CML;

KM blast crisis CML; BC-CML; human; chronic myelogenous leukaemia;

XX gene marker.

OS Homo sapiens.

XX US2003104426-A1.

XX 05-JUN-2003.

XX 14-JUN-2002; 2002US-00171561.

XX 18-JUN-2001; 2001US-0298914P.

XX (LINS/) LINSLEY P S.

XX (MAOM/) MAO M.

XX (DAIH/) DAI H.

XX (HEY/) HE Y.

XX (RAD/) RADICH J P.

XX Linsley PS, Mao M, Dai H, He Y, Radich JP,

XX WPI; 2003-787046/74.

XX Classifying cell sample as chronic phase chronic myelogenous leukemia or

XX blast crisis chronic myelogenous leukemia by detecting difference in

XX expression of genes corresponding to the markers such as X15415, U89436.

XX Disclosure; SEQ ID NO 9; 31pp; English.

XX The invention relates to a method of classifying a cell sample as chronic

XX phase chronic myelogenous leukemia (CP-CML) or blast crisis CML (BC-

XX CML). The method is useful for classifying a sample as CP-CML or BC-CML.

XX The present sequence represents a human chronic myelogenous leukemia

XX (CML) gene marker used to distinguish blast crisis CML from chronic phase

XX CML.

XX Sequence 1717 BP; 420 A; 343 C; 465 G; 489 T; 0 U; 0 Other;

XX Query Match 30.0%; Score 18; DB 10; Length 1717;

XX Best Local Similarity 100.0%; Pred. No. 5.6;

XX Matches 18; Conservative 0; Mismatches 0; Indels 0; Gaps 0;

QY 41 ACATGCGTGAAGCAGGTG 58

DB 515 ACATGCGTGAAGCAGGTG 532

RESULT 14

ID ADP07658

AC ADP07658 standard; DNA; 1926 BP.

DT 12-AUG-2004 (first entry)

DE Human secreted protein encoding DNA, seq id 141.

XX Cytostatic; antidiabetic; anorectic; gynaecological; antiparasitic;

KW dermatological; antiarteriosclerotic; antischismatic; neuroprotective;

KW neurotropic; antiparkinsonian; nephroprotective; human; secreted protein;

KW diagnostic; pharmaceutical; cancer; lung; oesophageal; liver; diabetes;

XX	obesity; metabolic disorder; cardiovascular disorder;
KW	reproductive disorder; psoriasis; eczema; bronchitis; cystic fibrosis;
KW	atherosclerosis; benign prostatic hyperplasia; asthma;
XX	Alzheimer's disease; Parkinson's disease; renal disorders; gene; ds.
OS	Homo sapiens.
XX	
XX	WO2004042000-A2.
XX	
XX	21-MAY-2004.
XX	
XX	16-MAY-2003; 2003WO-US015439.
XX	
XX	17-MAY-2002; 2002US-0381592P.
PR	12-JUN-2002; 2002US-0388543P.
PR	08-AUG-2002; 2002US-0401757P.
PR	12-AUG-2002; 2002US-0402585P.
PR	13-AUG-2002; 2002US-0402789P.
PR	22-AUG-2002; 2002US-0404592P.
PR	04-OCT-2002; 2002US-0415902P.
XX	
PA	(HUMA-) HUMAN GENOME SCI INC.
XX	
P1	Rosen CA, Ruben SM, Olsen H, Baker KP, Fiscella M, Wei P;
P1	Blrse CE, Komatsoulis G, Choi GH, Moore PA, Gupta R, Shi Y;
XX	
XX	WPI: 2004-400658/37.
DR	P-PSDB; ADP07840.
XX	
PT	New human secreted polypeptides and nucleic acid molecules for
PT	diagnosing, preventing or treating disorders associated with the secreted
PT	proteins, such as cancer, diabetes, obesity, cardiovascular disorders or
XX	renal disorders.
PS	Claim 7; SEQ ID NO 141; 1157bp; English.
XX	
CC	The invention relates to a human secreted polypeptide for diagnosing,
CC	preventing or treating disorders associated with the secreted proteins.
CC	The polypeptides and nucleic acid molecules of the invention are useful
CC	for preparing a diagnostic or pharmaceutical composition for diagnosing
CC	or treating a medical condition. These may be used for diagnosing,
CC	preventing or treating disorders related to the human secreted proteins,
CC	such as cancer (e.g. lung, oesophageal or liver cancer), diabetes,
CC	obesity, metabolic disorders, cardiovascular disorders, reproductive
CC	disorders, psoriasis, eczema, bronchitis, cystic fibrosis,
CC	atherosclerosis, benign prostatic hyperplasia, asthma, Alzheimer's
CC	disease, Parkinson's disease or renal disorders. Sequences given in
CC	records for ADP07558-ADP07709 represent human secreted protein encoding
XX	DNA's of the invention.
XX	
SO	Sequence 1926 BP; 479 A; 378 C; 568 G; 498 T; 0 U; 3 Other;
XX	
XX	Query Match 30.0%; Score 18; DB 12; Length 1926;
XX	Best Local Similarity 100.0%; Pred. NO. 5.6;
XX	Matches 18; Conservative 0; Mismatches 0; Indels 0; Gaps 0;
QY	41 ACATGCGTGAAGCAGGTG 58
DB	997 ACATGCGTGAAGCAGGTG 1014
XX	
XX	RESULT 15
XX	ADD22452
XX	ID ADD22452 standard; DNA; 2765 BP.
XX	ADD22452;
XX	15-JAN-2004 (first entry)
XX	
DE	H1A-Ba6 T cell recognised tumour antigenic polypeptide, SEQ No 102.
XX	
XX	tumour antigenic peptide; cancer; vaccine; cytostatic; cytotoxic T cell;
XX	colon; mouth; lung; prostatic; gynecological; human; gene; ds.

XX Homo sapiens.
 XX JP2003111595-A.
 XX 15-APR-2003.
 XX 24-JUN-2002; 2002JP-00183603.
 XX 25-JUN-2001; 2001JP-00191974.
 XX (ITOX/). ITO Y.
 XX WPI; 2003-611129/58.
 XX Novel tumor antigenic peptide or polypeptide useful for inducing
 XX cytotoxic T cells or for treating cancer such as colon, mouth, lung,
 XX prostate or gynecological cancer.
 XX Claim 10; SEQ ID NO 102; 98pp; Japanese.
 XX
 XX The invention relates to a novel tumour antigenic peptide or polypeptide
 XX comprising a sequence selected from 99 sequences fully defined in the
 XX specification. The tumour antigenic peptide or polypeptide comprises a
 XX sequence selected from 99 sequences fully defined in the specification,
 XX where the tumour antigenic peptide preferably has a sequence of Glu-Pro-
 XX Pro-Leu-Ser-Gln-Glu-Ile-Phe, and the polypeptide preferably has a
 XX sequence comprising 393 amino acids fully defined in the specification.
 XX The invention further provides a cancer vaccine comprising a tumour
 XX antigenic peptide or polypeptide, which has cytostatic activity. A tumour
 XX antigenic peptide, or polypeptide, its encoding polynucleotide, a
 XX hybridising polynucleotide, a recombinant vector containing the
 XX polynucleotide, a host transformed with the vector or an antibody are
 XX useful for screening for compounds that interact with the tumour
 XX antigenic peptide, the polypeptide or its encoding polynucleotide and
 XX increases the expression of the tumour antigenic peptide, the polypeptide
 XX or polynucleotide. The tumour antigenic peptide or the polypeptide is
 XX useful for inducing cytotoxic T cells. The tumour antigenic peptide
 XX vaccine is useful for treating cancer such as colon, mouth, lung,
 XX prostatic or gynecological cancer. The invention also provides a
 XX pharmaceutical composition useful for treating cancer. The tumour
 XX antigenic peptide or the polypeptide is useful as an antigen to create
 XX antibodies. This polynucleotide sequence represents the DNA encoding one
 XX of the human tumour antigenic polypeptides of the invention.
 XX
 XX Sequence 2765 BP; 759 A; 500 C; 652 G; 854 T; 0 U; 0 Other;
 XX
 XX Query Match 30.0%; Score 18; DB 10; Length 2765;
 XX Best Local Similarity 100.0%; Pred. No. 5.6;
 XX Matches 18; Conservative 0; Mismatches 0; Indels 0; Gaps 0.
 XX
 XX 41 ACATGGCTAAGCAGGTG 58
 XX |||||
 XX Db 509 ACATGGCTAAGCAGGTG 526
 XX |||||

Search completed: December 22, 2004, 22:44:12
 Job time : 217.015 secs

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OM nucleic - nucleic search, using sw model

Run on: December 22, 2004, 22:08:48 / Search time 48.3259 Seconds

(without alignments)
882.496 Million cell updates/sec

Title: US-09-898-616A-2

Perfect score: 60

Sequence: 1 agctacgaagcagctatgga.....acatgcgtgaagcagctgct 60

Scoring table: OLIGO NUC
Gapop 60.0, Gapext 60.0

Searched: 824507 seqs, 353394441 residues

Word size: 0

Total number of hits satisfying chosen parameters: 1649014

Minimum DB seq length: 0

Maximum DB seq length: 2000000000

Post-processing: Listing first 45 summaries

Database: Issued Patents, NA:

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6: /cgn2_6/ptodata/1/ina/backfile1.seq:*

Pred. No. is the number of results predicted by chance to have a score greater than or equal to the score of the result being printed, and is derived by analysis of the total score distribution.

SUMMARIES

Result No.	Score	Query Match	Length	ID	Description
1	60	100.0	60	US-08-864-357F-7	Sequence 7, Appl1
2	37	61.7	60	US-08-864-357F-12	Sequence 12, Appl1
3	18	30.0	59	US-08-864-357F-11	Sequence 11, Appl1
4	16	26.7	624	US-09-328-352-1928	Sequence 1928, Ap
5	16	26.7	830	US-09-023-655-504	Sequence 504, App
6	16	26.7	1959	US-09-252-991A-7983	Sequence 7983, Ap
7	16	26.7	2010	US-09-252-991A-8169	Sequence 8169, Ap
8	16	26.7	26700	US-08-472-217-1	Sequence 1, Appl1
9	16	26.7	26700	US-08-488-199-5	Sequence 5, Appl1
10	16	26.7	26700	US-08-760-534A-1	Sequence 1, Appl1
11	16	26.7	26700	US-09-335-757-1	Sequence 1, Appl1
12	15	25.0	540	US-09-270-767-3339	Sequence 3339, Ap
13	15	25.0	540	US-09-270-767-18621	Sequence 18621, A
14	15	25.0	1089	US-08-978-589A-1	Sequence 1, Appl1
15	15	25.0	1089	US-09-335-601-2	Sequence 2, Appl1
16	15	25.0	1089	US-09-219-120-1	Sequence 1, Appl1
17	15	25.0	1651	US-09-975-594-598	Sequence 598, App
18	15	25.0	1913	US-08-588-258A-41	Sequence 41, Appl1
19	15	25.0	1913	US-09-016-434-1078	Sequence 1078, Ap
20	15	25.0	1913	PCT-US98/08395-41	Sequence 41, Appl1
21	15	25.0	1946	US-08-785-584-1	Sequence 1, Appl1
22	15	25.0	1946	US-09-192-611-1	Sequence 1, Appl1
23	15	25.0	1946	US-08-755-592A-5	Sequence 5, Appl1
24	15	25.0	1946	US-09-617-923-1	Sequence 1, Appl1
25	15	25.0	2432	US-09-799-451-5	Sequence 5, Appl1
26	15	25.0	2513	US-09-799-451-6	Sequence 6, Appl1
27	15	25.0	2900	US-08-034-650-9	Sequence 9, Appl1

28	15	25.0	2900	US-08-449-015-9	Sequence 9, Appl1
29	15	25.0	6151	US-09-799-451-528	Sequence 528, App
30	15	25.0	7194	US-09-601-326-76	Sequence 76, Appl1
31	15	25.0	15420	US-09-601-326-54	Sequence 54, Appl1
32	14	23.3	136	US-09-513-999C-19445	Sequence 19445, A
33	14	23.3	292	US-09-313-294A-957	Sequence 957, App
34	14	23.3	303	US-09-489-039A-4877	Sequence 4877, Ap
35	14	23.3	363	US-09-248-786A-2610	Sequence 2610, Ap
36	14	23.3	373	US-09-513-999C-9517	Sequence 9517, Ap
37	14	23.3	498	US-09-621-976-2381	Sequence 2381, Ap
38	14	23.3	575	US-09-270-767-2759	Sequence 2759, A
39	14	23.3	724	US-08-998-416-810	Sequence 810, App
40	14	23.3	798	US-09-489-039A-4894	Sequence 4894, App
41	14	23.3	836	US-08-698-805-7	Sequence 7, Appl1
42	14	23.3	872	US-09-270-767-29234	Sequence 29234, A
43	14	23.3	879	US-09-620-312D-87	Sequence 87, Appl1
44	14	23.3	1105	US-09-221-017B-108	Sequence 108, App
45	14	23.3	1176	US-09-270-767-12926	Sequence 12926, A

ALIGNMENTS

RESULT 1
US-08-864-357F-7
Sequence 7, Application US/08864357F
Patent No. 6255281
GENERAL INFORMATION:
APPLICANT: Claragen, Inc. & NIH
TITLE OF INVENTION: Use of Recombinant Human Uteroglobulin in Treatment of Inflammato
FILE REFERENCE: 116142/2
CURRENT APPLICATION NUMBER: US/08/864,357F
CURRENT FILING DATE: 1997-05-28
NUMBER OF SEQ ID NOS: 22
SOFTWARE: Patentin version 3.0
SEQ ID NO 7
LENGTH: 60
TYPE: DNA
ORGANISM: artificial
FEATURE:
OTHER INFORMATION: primer sequence
US-08-864-357F-7
Query Match 100.0%; Score 60; DB 3; Length 60;
Best Local Similarity 100.0%; Pred. No. 2.5e-24;
Matches 60; Conservative 0; Mismatches 0; Indels 0; Gaps 0;
QY 1 AGCTACGAAGCAGCTATGGAAGCTTTCTTCGCGACGACATGCTGAAGCAGGTGCT 60
DB 1 AGCTACGAAGCAGCTATGGAAGCTTTCTTCGCGACGACATGCTGAAGCAGGTGCT 60
RESULT 2
US-08-864-357F-12/c
Sequence 12, Application US/08864357F
Patent No. 6255281
GENERAL INFORMATION:
APPLICANT: Claragen, Inc. & NIH
TITLE OF INVENTION: Use of Recombinant Human Uteroglobulin in Treatment of Inflammato
FILE REFERENCE: 116142/2
CURRENT APPLICATION NUMBER: US/08/864,357F
CURRENT FILING DATE: 1997-05-28
NUMBER OF SEQ ID NOS: 22
SOFTWARE: Patentin version 3.0
SEQ ID NO 12
LENGTH: 60
TYPE: DNA
ORGANISM: artificial
FEATURE:
OTHER INFORMATION: primer sequence
US-08-864-357F-12

Query Match 61.7%; Score 37; DB 3; Length 60;
Best Local Similarity 100.0%; Pred. No. 1.6e-11;
Matches 37; Conservative 0; Mismatches 0; Indels 0; Gaps 0;

QY 1 AGCTACGAGCAGCTATGAGACTGTTCTCCCGACC 37
DB 37 AGCTACGAGCAGCTATGAGACTGTTCTCCCGACC 1

RESULT 3
US-08-864-357F-11/C

Sequence 11, Application US/08864357F
Patent No. 6255281

GENERAL INFORMATION:

APPLICANT: Clontech, Inc. & NIH

TITLE OF INVENTION: Use of Recombinant Human Uteroglobin in Treatment of Inflammato

TITLE OF INVENTION: Fibrotic Conditions

FILE REFERENCE: 116142/2

CURRENT APPLICATION NUMBER: US/08/864,357F

CURRENT FILING DATE: 1997-05-28

NUMBER OF SEQ ID NOS: 22

SOFTWARE: Patent version 3.0

SEQ ID NO 11

LENGTH: 59

TYPE: DNA

ORGANISM: artificial

FEATURE:

OTHER INFORMATION: primer sequence

US-08-864-357F-11

Query Match 30.0%; Score 18; DB 3; Length 59;
Best Local Similarity 100.0%; Pred. No. 0.62;
Matches 18; Conservative 0; Mismatches 0; Indels 0; Gaps 0;

QY 38 AGGACATGCGTGAAGCAG 55
DB 59 AGGACATGCGTGAAGCAG 42

RESULT 4
US-09-328-352-1928

Sequence 1928, Application US/09328352

Patent No. 6562958

GENERAL INFORMATION:

APPLICANT: Gary L. Bretton et al.

TITLE OF INVENTION: NUCLEIC ACID AND AMINO ACID SEQUENCES RELATING TO ACINETOBACTER

FILE REFERENCE: GTC99-03BA

CURRENT APPLICATION NUMBER: US/09/328,352

CURRENT FILING DATE: 1999-06-04

NUMBER OF SEQ ID NOS: 8252

SEQ ID NO 1928

LENGTH: 624

TYPE: DNA

ORGANISM: Acinetobacter baumannii

US-09-328-352-1928

Query Match 26.7%; Score 16; DB 4; Length 624;
Best Local Similarity 100.0%; Pred. No. 8.3;
Matches 16; Conservative 0; Mismatches 0; Indels 0; Gaps 0;

QY 42 CATCGGTGAAGCAGT 57
DB 177 CATCGGTGAAGCAGT 192

RESULT 5
US-09-023-655-504/C

Sequence 504, Application US/09023655

Patent No. 6607879

GENERAL INFORMATION:

APPLICANT: Cocks, Benjamin G.

APPLICANT: Susan G. Stuart
APPLICANT: Jeffrey J. Seilhamer
TITLE OF INVENTION: COMPOSITION FOR THE DETECTION OF BLOOD CELL GENE
TITLE OF INVENTION: EXPRESSION
NUMBER OF SEQUENCES: 1508
CORRESPONDENCE ADDRESS:
ADDRESSEE: INCYTE PHARMACEUTICALS, INC.
STREET: 3174 PORTER DRIVE
CITY: PALO ALTO
STATE: CALIFORNIA
COUNTRY: USA
ZIP: 94304

COMPUTER READABLE FORM:

MEDIUM TYPE: Floppy disk

COMPUTER: IBM PC compatible

OPERATING SYSTEM: PC-DOS/MS-DOS

SOFTWARE: Word Perfect 6.1 for Windows/MS-DOS 6.2

CURRENT APPLICATION DATA:

APPLICATION NUMBER: US/09/023,655

FILING DATE: HEREMITH

CLASSIFICATION:

PRIOR APPLICATION DATA:

APPLICATION NUMBER:

FILING DATE:

CLASSIFICATION:

ATTORNEY/AGENT INFORMATION:

NAME: Zeller, Karen J.

REGISTRATION NUMBER: 37,071

REFERENCE/DOCKET NUMBER: PA-0001 US

TELEPHONE: (650) 855-0555

TELEFAX: (650) 845-4166

INFORMATION FOR SEQ ID NO: 504:

SEQUENCE CHARACTERISTICS:

LENGTH: 830 base pairs

TYPE: nucleic acid

STRANDEDNESS: single

TOPOLOGY: linear

IMMEDIATE SOURCE:

LIBRARY: TESTNOT03

CLONE: 2006402

US-09-023-655-504

Query Match 26.7%; Score 16; DB 4; Length 830;
Best Local Similarity 100.0%; Pred. No. 8.3;
Matches 16; Conservative 0; Mismatches 0; Indels 0; Gaps 0;

QY 45 GCGTGAAGCAGGTGCT 60
DB 195 GCGTGAAGCAGGTGCT 180

RESULT 6
US-09-252-991A-7983

Sequence 7983, Application US/09252991A

Patent No. 6551795

GENERAL INFORMATION:

APPLICANT: Marc J. Rubenfield et al.

TITLE OF INVENTION: NUCLEIC ACID AND AMINO ACID SEQUENCES RELATING TO PSEUDOMONAS

FILE REFERENCE: 107196.136

CURRENT APPLICATION NUMBER: US/09/252,991A

CURRENT FILING DATE: 1999-02-18

PRIOR APPLICATION NUMBER: US 60/074,788

PRIOR FILING DATE: 1998-02-18

PRIOR APPLICATION NUMBER: US 60/094,190

PRIOR FILING DATE: 1998-07-27

NUMBER OF SEQ ID NOS: 33142

SEQ ID NO 7983

LENGTH: 1959

TYPE: DNA

ORGANISM: Pseudomonas aeruginosa

US-09-252-991A-7983

Query Match 26.7%; Score 16; DB 4; Length 1959;
Best Local Similarity 100.0%; Pred. No. 8.4;
Matches 16; Conservative 0; Mismatches 0; Indels 0; Gaps 0;

QY 45 GCGTGAAGCAGGTGCT 60
DB 1913 GCGTGAAGCAGGTGCT 1928

RESULT 7
US-09-252-991A-8169/C
; Sequence 8169, Application US/09252991A
; Patent No. 6551795
; GENERAL INFORMATION:
; APPLICANT: Marc J. Rubenfield et al.
; TITLE OF INVENTION: NUCLEIC ACID AND AMINO ACID SEQUENCES RELATING TO PSEUDOMONAS
; FILE REFERENCE: 107196.136
; CURRENT APPLICATION NUMBER: US/09/252,991A
; PRIOR FILING DATE: 1999-02-18
; PRIOR APPLICATION NUMBER: US 60/074,788
; PRIOR FILING DATE: 1998-02-18
; PRIOR APPLICATION NUMBER: US 60/094,190
; PRIOR FILING DATE: 1998-07-27
; NUMBER OF SEQ ID NOS: 33142
; SEQ ID NO 8169
; LENGTH: 2010
; TYPE: DNA
; ORGANISM: Pseudomonas aeruginosa
US-09-252-991A-8169

Query Match 26.7%; Score 16; DB 4; Length 2010;
Best Local Similarity 100.0%; Pred. No. 8.4;
Matches 16; Conservative 0; Mismatches 0; Indels 0; Gaps 0;

QY 45 GCGTGAAGCAGGTGCT 60
DB 134 GCGTGAAGCAGGTGCT 119

RESULT 8
US-08-472-217-1
; Sequence 1, Application US/08472217
; Patent No. 5726058
; GENERAL INFORMATION:
; APPLICANT: Alanen-Kurki, Leena
; APPLICANT: Auvinen, Petri
; APPLICANT: Jaakkola, Panu
; APPLICANT: Jaakkola, Markku
; APPLICANT: Lepp, Sirpa
; APPLICANT: Maki, Markku
; APPLICANT: Vahinen, Tapani
; APPLICANT: Wrti, Anni
; TITLE OF INVENTION: Syndecan Stimulation Of Cellular
; TITLE OF INVENTION: Differentiation
; NUMBER OF SEQUENCES: 4
; CORRESPONDENCE ADDRESS:
; ADDRESSEE: Sterne, Kessler, Goldstein & Fox
; STREET: 1100 New York Avenue, Suite 600
; CITY: Washington
; STATE: D.C.
; COUNTRY: U.S.A.
; ZIP: 20005
; COMPUTER READABLE FORM:
; MEDIUM TYPE: Floppy disk
; COMPUTER: IBM PC compatible
; OPERATING SYSTEM: PC-DOS/MS-DOS
; SOFTWARE: Patentin Release #1.0, Version #1.25
; CURRENT APPLICATION DATA:
; APPLICATION NUMBER: US/08/472,217
; FILING DATE: 07-JUN-1995
; CLASSIFICATION: 514

PRIOR APPLICATION DATA:
; APPLICATION NUMBER: US 08/206,186
; FILING DATE: 07-MAR-1994
; PRIOR APPLICATION DATA:
; APPLICATION NUMBER: US 07/988,427
; FILING DATE: 01-DEC-1992
; ATTORNEY/AGENT INFORMATION:
; NAME: Cimbal, Michele A.
; REGISTRATION NUMBER: 33,851
; REFERENCE/DOCKET NUMBER: 1102,0050003
; TELECOMMUNICATION INFORMATION:
; TELEPHONE: (202) 371-2600
; TELEFAX: (202) 371-2540
; INFORMATION FOR SEQ ID NO: 1:
; SEQUENCE CHARACTERISTICS:
; LENGTH: 26700 base pairs
; TYPE: nucleic acid
; STRANDEDNESS: both
; TOPOLOGY: linear
; MOLECULE TYPE: DNA (genomic)
; HYPOTHETICAL: NO
; ANTI-SENSE: NO
; FEATURE:
; NAME/KEY: CDS
; LOCATION: join(4378..4443, 22026..22106, 23001..23483,
; ; LOCATION: 23905..24039, 24251..24418)
US-08-472-217-1

Query Match 26.7%; Score 16; DB 1; Length 26700;
Best Local Similarity 100.0%; Pred. No. 8.7;
Matches 16; Conservative 0; Mismatches 0; Indels 0; Gaps 0;

QY 29 CTCGGAGCAGACAT 44
DB 3800 CTCGGAGCAGACAT 3815

RESULT 9
US-08-488-199-5
; Sequence 5, Application US/08488199
; Patent No. 5851993
; GENERAL INFORMATION:
; APPLICANT: Jaakkola, Markku
; APPLICANT: Maki, Markku
; TITLE OF INVENTION: Suppression of Tumor Cell Growth By
; NUMBER OF SEQUENCES: 8
; CORRESPONDENCE ADDRESS:
; ADDRESSEE: STERNE, KESSLER, GOLDSTEIN & FOX
; STREET: 1100 New York Ave., NW
; CITY: Washington
; STATE: DC
; COUNTRY: USA
; ZIP: 20005
; COMPUTER READABLE FORM:
; MEDIUM TYPE: Floppy disk
; COMPUTER: IBM PC compatible
; OPERATING SYSTEM: PC-DOS/MS-DOS
; SOFTWARE: Patentin Release #1.0, Version #1.25
; CURRENT APPLICATION DATA:
; APPLICATION NUMBER: US/08/488,199
; FILING DATE: 07-JUN-1995
; CLASSIFICATION: 514
; PRIOR APPLICATION DATA:
; APPLICATION NUMBER: US 08/258,862
; FILING DATE: 13-JUN-1994
; ATTORNEY/AGENT INFORMATION:
; NAME: Cimbal, Michele A.
; REGISTRATION NUMBER: 33,851
; REFERENCE/DOCKET NUMBER: 1102,0130001
; TELECOMMUNICATION INFORMATION:
; TELEPHONE: 202-371-2600
; TELEFAX: 202-371-2540

INFORMATION FOR SEQ ID NO: 5:
SEQUENCE CHARACTERISTICS:
LENGTH: 26700 base pairs
TYPE: nucleic acid
STRANDEDNESS: single
TOPOLOGY: linear
FEATURE:
NAME/KEY: CDS
LOCATION: 4378..4443
FEATURE:
NAME/KEY: CDS
LOCATION: 22026..22107
FEATURE:
NAME/KEY: CDS
LOCATION: 23002..23483
FEATURE:
NAME/KEY: CDS
LOCATION: 23905..24040
FEATURE:
NAME/KEY: CDS
LOCATION: 24252..24418
US-08-488-199-5

Query Match: 26.7%; Score 16; DB 2; Length 26700;
Best Local Similarity 100.0%; Pred. No. 8.7;
Matches 16; Conservative 0; Mismatches 0; Indels 0; Gaps 0;

QY 29 CTCGGACCCAGGACAT 44
DB 3800 CTCGGACCCAGGACAT 3815

RESULT 10
US-08-760-534A-1
Sequence 1, Application US/08760534A
Patent No. 6017727
GENERAL INFORMATION:
APPLICANT: JALKANEN, MARKKU
APPLICANT: JAAKKOLA, PANU
APPLICANT: VIHINEN, TAPANI
TITLE OF INVENTION: SYNDECAN ENHANCER ELEMENT AND SYNDECAN
TITLE OF INVENTION: STIMULATION OF CELLULAR DIFFERENTIATION
NUMBER OF SEQUENCES: 14
CORRESPONDENCE ADDRESS:
ADDRESSEE: STERN, KESSLER, GOLDSTEIN & FOX P.L.L.C.
STREET: 1100 NEW YORK AVENUE, SUITE 600
CITY: WASHINGTON
STATE: DC
COUNTRY: US
ZIP: 20005-3934
COMPUTER READABLE FORM:
MEDIUM TYPE: floppy disk
COMPUTER: IBM PC compatible
OPERATING SYSTEM: PC-DOS/MS-DOS
SOFTWARE: Patentin Release #1.0, Version #1.30
CURRENT APPLICATION DATA:
APPLICATION NUMBER: US/08/760,534A
FILING DATE: 02-DEC-1996
CLASSIFICATION: 435
PRIORITY APPLICATION DATA:
APPLICATION NUMBER: US 08/206,186
FILING DATE: 07-MAR-1994
PRIOR APPLICATION DATA:
APPLICATION NUMBER: PCT/FI93/00514
FILING DATE: 01-DEC-1993
ATTORNEY/AGENT INFORMATION:
NAME: CIMBALA, MICHELE A.
REGISTRATION NUMBER: 33,851
REFERENCE/DOCKET NUMBER: 1708.0050004/MAC
TELECOMMUNICATION INFORMATION:
TELEPHONE: (202) 371-2600
TELEFAX: (202) 371-2540
INFORMATION FOR SEQ ID NO: 1:

SEQUENCE CHARACTERISTICS:
LENGTH: 26700 base pairs
TYPE: nucleic acid
STRANDEDNESS: double
TOPOLOGY: linear
MOLECULE TYPE: DNA (genomic)
FEATURE:
NAME/KEY: CDS
LOCATION: join(4378..4443, 22026..22106, 23001..23483,
23905..24039, 24251..24418)
US-08-760-534A-1

Query Match: 26.7%; Score 16; DB 3; Length 26700;
Best Local Similarity 100.0%; Pred. No. 8.7;
Matches 16; Conservative 0; Mismatches 0; Indels 0; Gaps 0;

QY 29 CTCGGACCCAGGACAT 44
DB 3800 CTCGGACCCAGGACAT 3815

RESULT 11
US-09-336-757-1
Sequence 1, Application US/09336757
Patent No. 6492344
GENERAL INFORMATION:
APPLICANT: JALKANEN, MARKKU
APPLICANT: JAAKKOLA, PANU
APPLICANT: VIHINEN, TAPANI
TITLE OF INVENTION: SYNDECAN ENHANCER ELEMENT AND SYNDECAN
TITLE OF INVENTION: STIMULATION OF CELLULAR DIFFERENTIATION
NUMBER OF SEQUENCES: 14
CORRESPONDENCE ADDRESS:
ADDRESSEE: STERN, KESSLER, GOLDSTEIN & FOX P.L.L.C.
STREET: 1100 NEW YORK AVENUE, SUITE 600
CITY: WASHINGTON
STATE: DC
COUNTRY: US
ZIP: 20005-3934
COMPUTER READABLE FORM:
MEDIUM TYPE: floppy disk
COMPUTER: IBM PC compatible
OPERATING SYSTEM: PC-DOS/MS-DOS
SOFTWARE: Patentin Release #1.0, Version #1.30
CURRENT APPLICATION DATA:
APPLICATION NUMBER: US/09/336,757
FILING DATE:
CLASSIFICATION:
PRIORITY APPLICATION DATA:
APPLICATION NUMBER: 08/760,534
FILING DATE:
PRIOR APPLICATION DATA:
APPLICATION NUMBER: PCT/FI93/00514
FILING DATE: 01-DEC-1993
ATTORNEY/AGENT INFORMATION:
NAME: CIMBALA, MICHELE A.
REGISTRATION NUMBER: 33,851
REFERENCE/DOCKET NUMBER: 1708.0050004/MAC
TELECOMMUNICATION INFORMATION:
TELEPHONE: (202) 371-2600
TELEFAX: (202) 371-2540
INFORMATION FOR SEQ ID NO: 1:
SEQUENCE CHARACTERISTICS:
LENGTH: 26700 base pairs
TYPE: nucleic acid
STRANDEDNESS: double
TOPOLOGY: linear
MOLECULE TYPE: DNA (genomic)
FEATURE:
NAME/KEY: CDS
LOCATION: join(4378..4443, 22026..22106, 23001..23483,
23905..24039, 24251..24418)
US-09-336-757-1

Query Match 26.7%; Score 16; DB 4; Length 26700;
Best Local Similarity 100.0%; Pred. No. 87;
Matches 16; Conservative 0; Mismatches 0; Indels 0; Gaps 0;

QY 29 CTCGACGACGACACT 44
DB 3800 CTCGACGACGACACT 3815

RESULT 12
US-09-270-767-3339
; Sequence 3339; Application US/09270767
; Patent No. 6703491
; GENERAL INFORMATION:
; APPLICANT: Homburger et al.
; TITLE OF INVENTION: Nucleic acids and proteins of *Drosophila melanogaster*
; FILE REFERENCE: File Reference: 7326-094
; CURRENT APPLICATION NUMBER: US/09/270,767
; CURRENT FILING DATE: 1999-03-17
; NUMBER OF SEQ ID NOS: 62517
; SOFTWARE: Patent In Ver. 2.0
; SEQ ID NO 3339
; LENGTH: 540
; TYPE: DNA
; ORGANISM: *Drosophila melanogaster*
; FEATURE:
; OTHER INFORMATION: n means any nucleotide
US-09-270-767-3339

Query Match 25.0%; Score 15; DB 4; Length 540;
Best Local Similarity 100.0%; Pred. No. 30;
Matches 15; Conservative 0; Mismatches 0; Indels 0; Gaps 0;

QY 10 GCAGCTATGGAAGTCTG 24
DB 134 GCAGCTATGGAAGTCTG 148

RESULT 13
US-09-270-767-18621
; Sequence 18621; Application US/09270767
; Patent No. 6703491
; GENERAL INFORMATION:
; APPLICANT: Homburger et al.
; TITLE OF INVENTION: Nucleic acids and proteins of *Drosophila melanogaster*
; FILE REFERENCE: File Reference: 7326-094
; CURRENT APPLICATION NUMBER: US/09/270,767
; CURRENT FILING DATE: 1999-03-17
; NUMBER OF SEQ ID NOS: 62517
; SOFTWARE: Patent In Ver. 2.0
; SEQ ID NO 18621
; LENGTH: 540
; TYPE: DNA
; ORGANISM: *Drosophila melanogaster*
; FEATURE:
; OTHER INFORMATION: n means any nucleotide
US-09-270-767-18621

Query Match 25.0%; Score 15; DB 4; Length 540;
Best Local Similarity 100.0%; Pred. No. 30;
Matches 15; Conservative 0; Mismatches 0; Indels 0; Gaps 0;

QY 10 GCAGCTATGGAAGTCTG 24
DB 134 GCAGCTATGGAAGTCTG 148

RESULT 14
US-08-978-589A-1
; Sequence 1; Application US/08978589A
; Patent No. 6087145
; GENERAL INFORMATION:

APPLICANT: ISHII, Takeshi
APPLICANT: MITSUDA, Satoshi
TITLE OF INVENTION: ESTERASE GENE AND ITS USE
NUMBER OF SEQUENCES: 4
CORRESPONDENCE ADDRESS:
ADDRESSEE: BIRCH, STEWART, KOLASCH & BIRCH, LLP
STREET: P.O. BOX 747
CITY: FALLS CHURCH
STATE: VIRGINIA
COUNTRY: UNITED STATES OF AMERICA
ZIP: 22040

COMPUTER READABLE FORM:
MEDIUM TYPE: Floppy disk
COMPUTER: IBM PC compatible
OPERATING SYSTEM: PC-DOS/MS-DOS
SOFTWARE: Patent In Release #1.0, Version #1.30
CURRENT APPLICATION DATA:
APPLICATION NUMBER: US/08/978,589A
FILING DATE: 26-NOV-1997

CLASSIFICATION: 435
ATTORNEY/AGENT INFORMATION:
NAME: Murphy Jr., Gerald M.
REGISTRATION NUMBER: 28,977
REFERENCE/DOCKET NUMBER: 20-4336P
TELECOMMUNICATION INFORMATION:
TELEPHONE: (703) 205-8000
TELEFAX: (703) 205-8050
INFORMATION FOR SEQ ID NO: 1:
SEQUENCE CHARACTERISTICS:
LENGTH: 1089 base pairs
TYPE: nucleic acid
STRANDEDNESS: double
TOPOLOGY: linear

MOLECULE TYPE: DNA (genomic)
ORGANISM: *Burkholderia cepacia*
STRAIN: SC-20
FEATURE:
NAME/KEY: CDS
LOCATION: 1..1089

US-08-978-589A-1

Query Match 25.0%; Score 15; DB 3; Length 1089;
Best Local Similarity 100.0%; Pred. No. 30;
Matches 15; Conservative 0; Mismatches 0; Indels 0; Gaps 0;

QY 46 CGTGAAGCAGGTGCT 60
DB 336 CGTGAAGCAGGTGCT 350

RESULT 15
US-09-336-601-2
; Sequence 2; Application US/09336601
; Patent No. 6184008
; GENERAL INFORMATION:
; APPLICANT: OHTA, Hiromichi
; APPLICANT: SUGAI, Takeshi
; APPLICANT: ISHII, Takeshi
; APPLICANT: MITSUDA, Satoshi
; TITLE OF INVENTION: PRODUCTION OF OPTICALLY ACTIVE SPHINGOID COMPOUND
; FILE REFERENCE: 2185-349P
; CURRENT APPLICATION NUMBER: US/09/336,601
; CURRENT FILING DATE: 1999-06-21
; EARLIER APPLICATION NUMBER: 09/034,007
; EARLIER FILING DATE: 1998-03-03
; NUMBER OF SEQ ID NOS: 2
; SOFTWARE: Patent In Ver. 2.0
; SEQ ID NO 2
; LENGTH: 1089
; TYPE: DNA
; ORGANISM: *E. coli* JM 109/pAL 612 strain
US-09-336-601-2

Query Match 25.0%; Score 15; DB 3; Length 1089;
 Best Local Similarity 100.0%; Pred. No. 30;
 Matches 15; Conservative 0; Mismatches 0; Indels 0; Gaps 0;

OY 46 CGTGAACGAGGTGCT 50
 |||||
 Db 336 CGTGAACGAGGTGCT 350

Search completed: December 23, 2004, 01:33:36
 Job time : 50.3258 secs

GenCore version 5.1.6
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OM nucleic - nucleic search, using sw model

Run on: December 22, 2004, 23:36:53 ; Search time 841.086 Seconds
(without alignments)
397.214 Million cell updates/sec

Title: US-09-898-616A-2

Perfect score: 60
Sequence: 1 agctacgaagcagctatgga.....acatgcgtgaagcagctgct 60

Scoring table: OLIGO_NUC
Gapop 60.0, Gapext 60.0

Searched: 410533 seqs, 2784055677 residues

Word size: 0

Total number of hits satisfying chosen parameters: 8210666

Minimum DB seq length: 0

Maximum DB seq length: 2000000000

Post-processing: Listing first 45 summaries

Database: Published Applications NA:*

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16: /cgn2_6/ptodata/1/pubna/US10D_PUBCOMB.seq:*
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19: /cgn2_6/ptodata/1/pubna/US11_NEW_PUB.seq:*
20: /cgn2_6/ptodata/1/pubna/US60_NEW_PUB.seq:*
21: /cgn2_6/ptodata/1/pubna/US60_PUBCOMB.seq:*

Pred. No. is the number of results predicted by chance to have a score greater than or equal to the score of the result being printed, and is derived by analysis of the total score distribution.

SUMMARIES

Result No.	Score	Query Match	Length DB	ID	Description
1	60	100.0	60	9 US-09-861-688-7	Sequence 7, Appl1
2	60	100.0	60	10 US-09-898-616A-2	Sequence 2, Appl1
3	60	100.0	60	15 US-10-187-498A-2	Sequence 2, Appl1
4	60	100.0	60	16 US-10-647-371-6	Sequence 2, Appl1
5	37	61.7	60	9 US-09-861-688-12	Sequence 12, Appl1
6	37	61.7	60	10 US-09-898-616A-7	Sequence 7, Appl1
7	37	61.7	60	10 US-09-898-616A-8	Sequence 8, Appl1
8	37	61.7	60	15 US-10-187-498A-7	Sequence 8, Appl1
9	37	61.7	60	15 US-10-187-498A-8	Sequence 8, Appl1
10	37	61.7	60	16 US-10-647-371-11	Sequence 11, Appl1
11	18	30.0	59	9 US-09-861-688-11	Sequence 11, Appl1
12	18	30.0	59	10 US-09-898-616A-6	Sequence 6, Appl1

ALIGNMENTS

c	13	18	30.0	59	15	US-10-187-498A-6	Sequence 6, Appl1
c	14	18	30.0	59	16	US-10-647-371-10	Sequence 10, Appl1
c	15	18	30.0	432	9	US-09-920-300A-1284	Sequence 1284, Ap
c	16	18	30.0	432	13	US-10-033-528-1284	Sequence 1284, Ap
c	17	18	30.0	432	15	US-10-099-926-1284	Sequence 1284, Ap
c	18	18	30.0	489	10	US-09-918-995-23144	Sequence 23144, A
c	19	18	30.0	1717	15	US-10-171-581-9	Sequence 9, Appl1
c	20	18	30.0	3252	18	US-10-723-860-5226	Sequence 5226, Ap
c	21	18	30.0	3299	15	US-10-006-285-513	Sequence 513, Appl1
c	22	17	28.3	467	13	US-10-027-632-59298	Sequence 59298, A
c	23	17	28.3	467	13	US-10-027-632-59300	Sequence 59300, A
c	24	17	28.3	467	13	US-10-027-632-59300	Sequence 59300, A
c	25	17	28.3	467	13	US-10-027-632-298628	Sequence 298628, A
c	26	17	28.3	467	13	US-10-027-632-298630	Sequence 298630, A
c	27	17	28.3	467	13	US-10-027-632-298630	Sequence 298630, A
c	28	17	28.3	467	15	US-10-027-632-59298	Sequence 59298, A
c	29	17	28.3	467	15	US-10-027-632-59299	Sequence 59299, A
c	30	17	28.3	467	15	US-10-027-632-59300	Sequence 59300, A
c	31	17	28.3	467	15	US-10-027-632-298628	Sequence 298628, A
c	32	17	28.3	467	15	US-10-027-632-298629	Sequence 298629, A
c	33	17	28.3	467	15	US-10-027-632-298630	Sequence 298630, A
c	34	17	28.3	534	13	US-10-027-632-50093	Sequence 50093, A
c	35	17	28.3	534	15	US-10-027-632-50093	Sequence 50093, A
c	36	17	28.3	555	13	US-10-027-632-321780	Sequence 321780, A
c	37	17	28.3	555	15	US-10-027-632-321780	Sequence 321780, A
c	38	16	26.7	65	10	US-09-908-975-28883	Sequence 28883, A
c	39	16	26.7	351	17	US-10-430-201-2182	Sequence 2182, Ap
c	40	16	26.7	373	17	US-10-430-201-2183	Sequence 2183, Ap
c	41	16	26.7	373	9	US-09-983-965-739	Sequence 739, Appl
c	42	16	26.7	375	18	US-10-674-124A-22078	Sequence 22078, A
c	43	16	26.7	477	9	US-09-783-590-4830	Sequence 4830, Ap
c	44	16	26.7	492	18	US-10-425-115-102427	Sequence 102427, A
c	45	16	26.7	561	13	US-10-027-632-10327	Sequence 10327, A

RESULT 1

US-09-861-688-7

Sequence 7, Application US/09861688

Patent No. US20020173460A1

GENERAL INFORMATION:

APPLICANT: Chazgen, Inc. & NIH

TITLE OF INVENTION: Use of Recombinant Human Uteroglobin in Treatment of

TITLE OF INVENTION: Inflammatory and

FILE REFERENCE: 116142/2

CURRENT APPLICATION NUMBER: US/09/861,688

CURRENT FILING DATE: 2001-05-21

PRIOR APPLICATION NUMBER: 09/864,357

PRIOR FILING DATE: 1997-05-28

NUMBER OF SEQ ID NOS: 22

SOFTWARE: PatentIn version 3.0

SEQ ID NO 7

LENGTH: 60

TYPE: DNA

ORGANISM: Artificial Sequence

FEATURE:

OTHER INFORMATION: primer sequence

US-09-861-688-7

Query Match 100.0%; Score 60; DB 9; Length 60;

Best Local Similarity 100.0%; Pred. No. 6,3e-24;

Matches 60; Conservative 0; Mismatches 0; Gaps 0;

DB 1 AGCTACGAAGCAGCTATGGAAGCTCTCTCCGACGACATGCGTGAAGAGAGTGTCT 60

1 AGCTACGAAGCAGCTATGGAAGCTCTCTCTCCGACGACATGCGTGAAGAGAGTGTCT 60

RESULT 2

US-09-898-616A-2

Sequence 2, Application US/09898616A
Publication No. US20030109429A1
GENERAL INFORMATION:
APPLICANT: Clargen Inc.
APPLICANT: Pilon, Aprile L
APPLICANT: Welch, Richard W
TITLE OF INVENTION: Method for the Production of Purified rhug for the Treatment of
TITLE OF INVENTION: Inflammatory and Fibrotic Conditions
FILE REFERENCE: 116142/00170
CURRENT APPLICATION NUMBER: US/09/898,616A
CURRENT FILING DATE: 2002-10-15
PRIOR APPLICATION NUMBER: US 08/864,357
PRIOR FILING DATE: 1997-05-28
NUMBER OF SEQ ID NOS: 10
SOFTWARE: PatentIn version 3.1
SEQ ID NO 2
LENGTH: 60
TYPE: DNA
ORGANISM: Artificial Sequence
FEATURE:
OTHER INFORMATION: Original amino acid sequence obtained from cDNA. Current submitte
OTHER INFORMATION: d sequence maximized for expression in E. coli.
NAME/KEY: misc feature
OTHER INFORMATION: Original amino acid sequence obtained from cDNA. Current submitte
OTHER INFORMATION: d sequence maximized for expression in E. coli.
US-09-898-616A-2

Query Match 100.0%; Score 60; DB 10; Length 60;
Best Local Similarity 100.0%; Pred. No. 6.3e-24;
Matches 60; Conservative 0; Mismatches 0; Indels 0; Gaps 0;

Db 1 AGCTACGAGAGGATGAGTCTCTCCGACGACGACATGCGTGAAGCAGGTGCT 60
1 AGCTACGAGAGGATGAGTCTCTCCGACGACGACATGCGTGAAGCAGGTGCT 60

RESULT 3
US-10-187-498A-2
Sequence 2, Application US/10187498A
Publication No. US2003020795A1
GENERAL INFORMATION:
APPLICANT: Clargen Inc.
APPLICANT: Pilon, Aprile L
APPLICANT: Welch, Richard W
TITLE OF INVENTION: Method for the Production of Purified rhug for the Treatment of
TITLE OF INVENTION: Inflammatory and Fibrotic Conditions
FILE REFERENCE: 116142/00260
CURRENT APPLICATION NUMBER: US/10/187,498A
CURRENT FILING DATE: 2001-07-02
PRIOR APPLICATION NUMBER: US 08/864,357
PRIOR FILING DATE: 1997-05-28
NUMBER OF SEQ ID NOS: 10
SOFTWARE: PatentIn version 3.1
SEQ ID NO 2
LENGTH: 60
TYPE: DNA
ORGANISM: Artificial Sequence
FEATURE:
OTHER INFORMATION: Original amino acid sequence obtained from cDNA. Current submitte
OTHER INFORMATION: d sequence maximized for expression in E. coli.
NAME/KEY: misc feature
OTHER INFORMATION: Original amino acid sequence obtained from cDNA. Current submitte
OTHER INFORMATION: d sequence maximized for expression in E. coli.
US-10-187-498A-2

Query Match 100.0%; Score 60; DB 15; Length 60;
Best Local Similarity 100.0%; Pred. No. 6.3e-24;
Matches 60; Conservative 0; Mismatches 0; Indels 0; Gaps 0;

Db 1 AGCTACGAGAGGATGAGTCTCTCCGACGACGACATGCGTGAAGCAGGTGCT 60
1 AGCTACGAGAGGATGAGTCTCTCCGACGACGACATGCGTGAAGCAGGTGCT 60

Db 1 AGCTACGAGAGGATGAGTCTCTCTCCGACGACGACATGCGTGAAGCAGGTGCT 60
1 AGCTACGAGAGGATGAGTCTCTCTCCGACGACGACATGCGTGAAGCAGGTGCT 60

RESULT 4
US-10-647-371-6
Sequence 6, Application US/10647371
Publication No. US20040047857A1
GENERAL INFORMATION:
APPLICANT: Clargen, Inc. & NIH
TITLE OF INVENTION: Use of Recombinant Human Uteroglobulin in Treatment of Inflammatory
TITLE OF INVENTION: and Fibrotic Conditions
FILE REFERENCE: 116142-85
CURRENT APPLICATION NUMBER: US/10/647,371
CURRENT FILING DATE: 2003-08-25
PRIOR APPLICATION NUMBER: 09/549,926
PRIOR FILING DATE: 2000-04-14
NUMBER OF SEQ ID NOS: 12
SOFTWARE: PatentIn version 3.2
SEQ ID NO 6
LENGTH: 60
TYPE: DNA
ORGANISM: Artificial
FEATURE:
OTHER INFORMATION: Description of Artificial Sequence: primer sequence
US-10-647-371-6

Query Match 100.0%; Score 60; DB 15; Length 60;
Best Local Similarity 100.0%; Pred. No. 6.3e-24;
Matches 60; Conservative 0; Mismatches 0; Indels 0; Gaps 0;

Db 1 AGCTACGAGAGGATGAGTCTCTCTCCGACGACGACATGCGTGAAGCAGGTGCT 60
1 AGCTACGAGAGGATGAGTCTCTCTCCGACGACGACATGCGTGAAGCAGGTGCT 60

RESULT 5
US-09-861-688-12/c
Sequence 12, Application US/09861688
Patent No. US20020173460A1
GENERAL INFORMATION:
APPLICANT: Clargen, Inc. & NIH
TITLE OF INVENTION: Use of Recombinant Human Uteroglobulin in Treatment of
TITLE OF INVENTION: Inflammatory and
FILE REFERENCE: 116142/2
CURRENT APPLICATION NUMBER: US/09/861,688
CURRENT FILING DATE: 2001-05-21
PRIOR APPLICATION NUMBER: 08/864,357
PRIOR FILING DATE: 1997-05-28
NUMBER OF SEQ ID NOS: 22
SOFTWARE: PatentIn version 3.0
SEQ ID NO 12
LENGTH: 60
TYPE: DNA
ORGANISM: Artificial Sequence
FEATURE:
OTHER INFORMATION: primer sequence
US-09-861-688-12

Query Match 61.7%; Score 37; DB 9; Length 60;
Best Local Similarity 100.0%; Pred. No. 6.3e-11;
Matches 37; Conservative 0; Mismatches 0; Indels 0; Gaps 0;

Db 1 AGCTACGAGAGGATGAGTCTCTCTCCGACGACGACATGCGTGAAGCAGGTGCT 37
37 AGCTACGAGAGGATGAGTCTCTCTCCGACGACGACATGCGTGAAGCAGGTGCT 1

RESULT 6
US-09-898-616A-7/c
Sequence 7, Application US/09898616A
Publication No. US20030109429A1
GENERAL INFORMATION:

APPLICANT: Clargen Inc.
APPLICANT: Pilon, Aprile L
TITLE OF INVENTION: Method for the Production of Purified rhug for the Treatment of
TITLE OF INVENTION: Inflammation and Fibrotic Conditions
FILE REFERENCE: 116142/00170
CURRENT APPLICATION NUMBER: US/09/898,616A
PRIOR FILING DATE: 2002-10-15
PRIOR APPLICATION NUMBER: US 08/864,357
PRIOR FILING DATE: 1997-05-28
NUMBER OF SEQ ID NOS: 10
SOFTWARE: PatentIn version 3.1
SEQ ID NO 7
LENGTH: 60
TYPE: DNA
ORGANISM: Artificial Sequence
FEATURE:
OTHER INFORMATION: Original amino acid sequence obtained from cDNA. Current submitte
FEATURE:
OTHER INFORMATION: d sequence maximized for expression in E. coli.
NAME/KEY: misc feature
OTHER INFORMATION: Original amino acid sequence obtained from cDNA. Current submitte
OTHER INFORMATION: d sequence maximized for expression in E. coli.
US-09-898-616A-7

Query Match 61.7%; Score 37; DB 10; Length 60;
Best Local Similarity 100.0%; Pred. No. 6.3e-11;
Matches 37; Conservative 0; Mismatches 0; Gaps 0;

QY 1 AGCTACGAGCAGCTATGGAAGTCTCTCCGAGCC 37
DB 37 AGCTACGAGCAGCTATGGAAGTCTCTCCGAGCC 1

RESULT 7
US-09-898-616A-8/c
Sequence 8, Application US/09898616A
Publication No. US20030109429A1
GENERAL INFORMATION:
APPLICANT: Clargen Inc.
APPLICANT: Pilon, Aprile L
TITLE OF INVENTION: Method for the Production of Purified rhug for the Treatment of
TITLE OF INVENTION: Inflammation and Fibrotic Conditions
FILE REFERENCE: 116142/00170
CURRENT APPLICATION NUMBER: US/09/898,616A
PRIOR FILING DATE: 2002-10-15
PRIOR APPLICATION NUMBER: US 08/864,357
PRIOR FILING DATE: 1997-05-28
NUMBER OF SEQ ID NOS: 10
SOFTWARE: PatentIn version 3.1
SEQ ID NO 8
LENGTH: 60
TYPE: DNA
ORGANISM: Artificial Sequence
FEATURE:
OTHER INFORMATION: Original amino acid sequence obtained from cDNA. Current submitte
FEATURE:
OTHER INFORMATION: d sequence maximized for expression in E. coli.
NAME/KEY: misc feature
OTHER INFORMATION: Original amino acid sequence obtained from cDNA. Current submitte
OTHER INFORMATION: d sequence maximized for expression in E. coli.
US-09-898-616A-8

Query Match 61.7%; Score 37; DB 10; Length 60;
Best Local Similarity 100.0%; Pred. No. 6.3e-11;
Matches 37; Conservative 0; Mismatches 0; Indels 0; Gaps 0;

QY 1 AGCTACGAGCAGCTATGGAAGTCTCTCCGAGCC 37
DB 37 AGCTACGAGCAGCTATGGAAGTCTCTCCGAGCC 1

RESULT 8
US-10-187-498A-7/c
Sequence 7, Application US/10187498A
Publication No. US20030207795A1
GENERAL INFORMATION:
APPLICANT: Clargen Inc.
APPLICANT: Pilon, Aprile L
TITLE OF INVENTION: Method for the Production of Purified rhug for the Treatment of
TITLE OF INVENTION: Inflammation and Fibrotic Conditions
FILE REFERENCE: 116142/00260
CURRENT APPLICATION NUMBER: US/10/187,498A
PRIOR FILING DATE: 2001-07-02
PRIOR APPLICATION NUMBER: US 08/864,357
PRIOR FILING DATE: 1997-05-28
NUMBER OF SEQ ID NOS: 10
SOFTWARE: PatentIn version 3.1
SEQ ID NO 7
LENGTH: 60
TYPE: DNA
ORGANISM: Artificial Sequence
FEATURE:
OTHER INFORMATION: Original amino acid sequence obtained from cDNA. Current submitte
FEATURE:
OTHER INFORMATION: d sequence maximized for expression in E. coli.
NAME/KEY: misc feature
OTHER INFORMATION: Original amino acid sequence obtained from cDNA. Current submitte
OTHER INFORMATION: d sequence maximized for expression in E. coli.
US-10-187-498A-7

Query Match 61.7%; Score 37; DB 15; Length 60;
Best Local Similarity 100.0%; Pred. No. 6.3e-11;
Matches 37; Conservative 0; Mismatches 0; Indels 0; Gaps 0;

QY 1 AGCTACGAGCAGCTATGGAAGTCTCTCCGAGCC 37
DB 37 AGCTACGAGCAGCTATGGAAGTCTCTCCGAGCC 1

RESULT 9
US-10-187-498A-8/c
Sequence 8, Application US/10187498A
Publication No. US20030207795A1
GENERAL INFORMATION:
APPLICANT: Clargen Inc.
APPLICANT: Pilon, Aprile L
TITLE OF INVENTION: Method for the Production of Purified rhug for the Treatment of
TITLE OF INVENTION: Inflammation and Fibrotic Conditions
FILE REFERENCE: 116142/00260
CURRENT APPLICATION NUMBER: US/10/187,498A
PRIOR FILING DATE: 2001-07-02
PRIOR APPLICATION NUMBER: US 08/864,357
PRIOR FILING DATE: 1997-05-28
NUMBER OF SEQ ID NOS: 10
SOFTWARE: PatentIn version 3.1
SEQ ID NO 8
LENGTH: 60
TYPE: DNA
ORGANISM: Artificial Sequence
FEATURE:
OTHER INFORMATION: Original amino acid sequence obtained from cDNA. Current submitte
FEATURE:
OTHER INFORMATION: d sequence maximized for expression in E. coli.
NAME/KEY: misc feature
OTHER INFORMATION: Original amino acid sequence obtained from cDNA. Current submitte
OTHER INFORMATION: d sequence maximized for expression in E. coli.
US-10-187-498A-8

Query Match 61.7%; Score 37; DB 15; Length 60;
Best Local Similarity 100.0%; Pred. No. 6.3e-11;
Matches 37; Conservative 0; Mismatches 0; Indels 0; Gaps 0;

OY 1 AGTACGAGAGAGCTATGAGTCTCTCTCCGACC 37
DB 37 AGTACGAGAGAGCTATGAGTCTCTCTCCGACC 1

RESULT 10
US-10-647-371-11/c
Sequence 11, Application US/10647371
Publication No. US20040047857A1
GENERAL INFORMATION:
APPLICANT: Claragen, Inc. & NIH
TITLE OF INVENTION: Use of Recombinant Human Uteroglobulin in Treatment of Inflammatory
TITLE OF INVENTION: and Fibrotic Conditions
FILE REFERENCE: 116142-85
CURRENT APPLICATION NUMBER: US/10/647,371
CURRENT FILING DATE: 2003-08-25
PRIOR APPLICATION NUMBER: 09/549,926
PRIOR FILING DATE: 2000-04-14
NUMBER OF SEQ ID NOS: 12
SOFTWARE: PatentIn version 3.2
SEQ ID NO 11
LENGTH: 60
TYPE: DNA
ORGANISM: Artificial
FEATURE:
OTHER INFORMATION: Description of Artificial Sequence: primer sequence
US-10-647-371-11

Query Match 61.7%; Score 37; DB 16; Length 60;
Best Local Similarity 100.0%; Pred. No. 6,3e-11;
Matches 37; Conservative 0; Mismatches 0; Indels 0; Gaps 0;

OY 1 AGTACGAGAGAGCTATGAGTCTCTCTCCGACC 37
DB 37 AGTACGAGAGAGCTATGAGTCTCTCTCCGACC 1

RESULT 11
US-09-861-688-11/c
Sequence 11, Application US/09861688
Patent No. US20020173460A1
GENERAL INFORMATION:
APPLICANT: Claragen, Inc. & NIH
TITLE OF INVENTION: Use of Recombinant Human Uteroglobulin in Treatment of
TITLE OF INVENTION: Inflammatory and
TITLE OF INVENTION: Fibrotic Conditions
FILE REFERENCE: 116142/2
CURRENT APPLICATION NUMBER: US/09/861,688
CURRENT FILING DATE: 2001-05-21
PRIOR APPLICATION NUMBER: 08/864,357
PRIOR FILING DATE: 1997-05-28
NUMBER OF SEQ ID NOS: 22
SOFTWARE: PatentIn version 3.0
SEQ ID NO 11
LENGTH: 59
TYPE: DNA
ORGANISM: Artificial Sequence
FEATURE:
OTHER INFORMATION: primer sequence
US-09-861-688-11

Query Match 30.0%; Score 18; DB 9; Length 59;
Best Local Similarity 100.0%; Pred. No. 3.4;
Matches 18; Conservative 0; Mismatches 0; Indels 0; Gaps 0;

OY 38 AGGACATGCGTGAAGCAG 55
DB 59 AGGACATGCGTGAAGCAG 42

RESULT 12
US-09-898-616a-6/c
Sequence 6, Application US/09898616A

Publication No. US20030109429A1
GENERAL INFORMATION:
APPLICANT: Claragen Inc.
APPLICANT: Pilon, Aprile L
APPLICANT: Welch, Richard W
TITLE OF INVENTION: Method for the Production of Purified rhug for the Treatment of
TITLE OF INVENTION: Inflammatory and Fibrotic Conditions
FILE REFERENCE: 116142/00170
CURRENT APPLICATION NUMBER: US/09/898,616A
CURRENT FILING DATE: 2002-10-15
PRIOR APPLICATION NUMBER: US 08/864,357
PRIOR FILING DATE: 1997-05-28
NUMBER OF SEQ ID NOS: 10
SOFTWARE: PatentIn version 3.1
SEQ ID NO 6
LENGTH: 59
TYPE: DNA
ORGANISM: Artificial Sequence
FEATURE:
OTHER INFORMATION: Original amino acid sequence obtained from cDNA. Current submitte
OTHER INFORMATION: d sequence maximized for expression in E. coli.
FEATURE:
NAME/KEY: misc. feature
OTHER INFORMATION: Original amino acid sequence obtained from cDNA. Current submitte
OTHER INFORMATION: d sequence maximized for expression in E. coli.
US-09-898-616a-6

Query Match 30.0%; Score 18; DB 10; Length 59;
Best Local Similarity 100.0%; Pred. No. 3.4;
Matches 18; Conservative 0; Mismatches 0; Indels 0; Gaps 0;

OY 38 AGGACATGCGTGAAGCAG 55
DB 59 AGGACATGCGTGAAGCAG 42

RESULT 13
US-10-187-498a-6/c
Sequence 6, Application US/10187498A
Publication No. US20030207795A1
GENERAL INFORMATION:
APPLICANT: Claragen Inc.
APPLICANT: Pilon, Aprile L
APPLICANT: Welch, Richard W
TITLE OF INVENTION: Method for the Production of Purified rhug for the Treatment of
TITLE OF INVENTION: Inflammatory and Fibrotic Conditions
FILE REFERENCE: 116142/00260
CURRENT APPLICATION NUMBER: US/10/187,498A
CURRENT FILING DATE: 2001-07-02
PRIOR APPLICATION NUMBER: US 08/864,357
PRIOR FILING DATE: 1997-05-28
NUMBER OF SEQ ID NOS: 10
SOFTWARE: PatentIn version 3.1
SEQ ID NO 6
LENGTH: 59
TYPE: DNA
ORGANISM: Artificial Sequence
FEATURE:
OTHER INFORMATION: Original amino acid sequence obtained from cDNA. Current submitte
OTHER INFORMATION: d sequence maximized for expression in E. coli.
FEATURE:
NAME/KEY: misc. feature
OTHER INFORMATION: Original amino acid sequence obtained from cDNA. Current submitte
OTHER INFORMATION: d sequence maximized for expression in E. coli.
US-10-187-498a-6

Query Match 30.0%; Score 18; DB 15; Length 59;
Best Local Similarity 100.0%; Pred. No. 3.4;
Matches 18; Conservative 0; Mismatches 0; Indels 0; Gaps 0;

OY 38 AGGACATGCGTGAAGCAG 55
DB 59 AGGACATGCGTGAAGCAG 42

RESULT 14

US-10-647-371-10/C
; Sequence 10, Application US/10647371
; Publication No. US20040047857A1
; GENERAL INFORMATION:
; APPLICANT: Clargen, Inc. & NIH
; TITLE OF INVENTION: Use of Recombinant Human Uteroglobulin in Treatment of Inflammatory
; TITLE OF INVENTION: and Fibrotic Conditions
; FILE REFERENCE: 116142-85
; CURRENT APPLICATION NUMBER: US/10/647,371
; CURRENT FILING DATE: 2003-08-25
; PRIOR APPLICATION NUMBER: 09/549,926
; PRIOR FILING DATE: 2000-04-14
; NUMBER OF SEQ ID NOS: 12
; SOFTWARE: PatentIn version 3.2
; SEQ ID NO 10
; LENGTH: 59
; TYPE: DNA
; ORGANISM: Artificial
; FEATURE:
; OTHER INFORMATION: Description of Artificial Sequence: primer sequence
US-10-647-371-10

Query Match

30.0%; Score 18; DB 16; Length 59;
Best Local Similarity 100.0%; Pred. No. 3.4;
Matches 18; Conservative 0; Mismatches 0; Indels 0; Gaps 0;

QY 38 AGGACATGCGTGAAGCAG 55
|||
Db 59 AGGACATGCGTGAAGCAG 42

RESULT 15

US-09-920-300A-1284
; Sequence 1284, Application US/09920300A
; Patent No. US20020136728A1
; GENERAL INFORMATION:
; APPLICANT: King, Gordon E.
; APPLICANT: Meagher, Madeleine Joy
; APPLICANT: Xu, Jiangchun
; APPLICANT: Secrist, Heather
; TITLE OF INVENTION: COMPOSITIONS AND METHODS FOR THE THERAPY
; TITLE OF INVENTION: AND DIAGNOSIS OF COLON CANCER
; FILE REFERENCE: 210121.547
; CURRENT APPLICATION NUMBER: US/09/920,300A
; CURRENT FILING DATE: 2001-07-31
; NUMBER OF SEQ ID NOS: 1789
; SOFTWARE: FastSeq for Windows Version 4.0
; SEQ ID NO 1284
; LENGTH: 432
; TYPE: DNA
; ORGANISM: Homo sapiens
US-09-920-300A-1284

Query Match

30.0%; Score 18; DB 9; Length 432;
Best Local Similarity 100.0%; Pred. No. 3;
Matches 18; Conservative 0; Mismatches 0; Indels 0; Gaps 0;

QY 41 ACATGCGTGAAGCAGGTG 58
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Db 9 ACATGCGTGAAGCAGGTG 26

Search completed: December 23, 2004, 05:19:26
Job time : 843.086 secs

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OM nucleic - nucleic search, using sw model

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(without alignments)
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Title: US-09-898-616A-3

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Gapop 60.0, Gapext 60.0

Scoring table: 4525729 seqs, 2364849745 residues

Searched: 0

Word size: 0

Total number of hits satisfying chosen parameters: 9053458

Minimum DB seq length: 0

Maximum DB seq length: 200000000

Post-processing: listing first 45 summaries

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2: gb_htg.*
3: gb_in.*
4: gb_cm.*
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9: gb_pr.*
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11: gb_scs.*
12: gb_sy.*
13: gb_un.*
14: gb_vi.*

Pred. No. is the number of results predicted by chance to have a score greater than or equal to the score of the result being printed, and is derived by analysis of the total score distribution.

SUMMARIES

Result No.	Score	Query Match	Length DB	ID	Description
1	59	100.0	59	6	ARI60917 Sequence
2	37	62.7	59	6	ARI60920 Sequence
3	19	32.2	62318	2	ACI35267 Rattus no
4	19	32.2	65394	2	ACI17645 Mus muscu
5	19	32.2	145825	8	OSJN00027
6	19	32.2	232344	2	ACI11782 Rattus no
7	19	32.2	246787	2	ACI28608 Rattus no
8	18	30.5	215	6	AI3577
9	18	30.5	215	6	AI3577
10	18	30.5	220	12	SYNBILRG
11	18	30.5	1234	8	AX461306 Sequence
12	18	30.5	2004	6	AX461306 Sequence
13	18	30.5	3176	8	AX065797 Oryza sat
14	18	30.5	7419	1	PSETPEDC
15	18	30.5	42499	8	AC004625 Arabidops
16	18	30.5	67388	9	ACI33476 Homo sapi
17	18	30.5	96066	8	AC005662 Arabidops
18	18	30.5	106057	8	ACI19796 Oryza sat
19	18	30.5	110000	1	RME591985_15

c 20	18	30.5	150503	8	ACI22149 Oryza sat
c 21	18	30.5	159867	2	ACI30411 Homo sapi
c 22	18	30.5	166399	2	AC012584 Homo sapi
c 23	18	30.5	169199	2	ACI08292 Mus muscu
c 24	18	30.5	170064	8	AF003771 Oryza sat
c 25	18	30.5	193386	10	ACI17693 Arabidops
c 26	18	30.5	202920	10	AL929441 Mouse DNA
c 27	18	30.5	207925	2	ACI26948 Rattus no
c 28	17	28.8	195	6	AX435021
c 29	17	28.8	993	13	AY487489
c 30	17	28.8	2190	10	AF144255
c 31	17	28.8	3530	6	CO582203
c 32	17	28.8	5568	6	AX571642
c 33	17	28.8	10166	1	AE007324 Streptoco
c 34	17	28.8	11627	1	AE012368 Xanthomon
c 35	17	28.8	14736	6	CQ789080 Sequence
c 36	17	28.8	14736	6	AR218939 Sequence
c 37	17	28.8	14736	6	BD003851 Polynucle
c 38	17	28.8	19457	2	AC017722 Drosophila
c 39	17	28.8	62598	6	AX571767 Sequence
c 40	17	28.8	65904	2	ACI04954 Homo sapi
c 41	17	28.8	91803	9	AC092318 Homo sapi
c 42	17	28.8	95506	9	AC002407 Human Chr
c 43	17	28.8	100061	2	AC016551 Homo sapi
c 44	17	28.8	132164	9	AC098587 Homo sapi
c 45	17	28.8	144402	2	AC021854 Homo sapi

ALIGNMENTS

RESULT 1	ARI60917	59 bp	DNA	linear	PAT 17-OCT-2001
LOCUS	ARI60917	Sequence 8 from patent US 6255281.			
DEFINITION	ARI60917				
ACCESSION	ARI60917.1	GI:16225984			
VERSION					
KEYWORDS					
SOURCE	Unknown.				
ORGANISM	Unknown.				
REFERENCE	1 (bases 1 to 59)				
AUTHORS	Pilon A.L., Mukherjee, A.B. and Zhang, Z.				
TITLE	Use of recombinant human uteroglobin in treatment of inflammatory				
JOURNAL	Patent: US 6255281-A 8 03-JUL-2001;				
FEATURES	Location/Qualifiers				
source	1..59				
ORIGIN	/organism="unknown"				
	/mol_type="unassigned DNA"				
Query Match	100.0%;	Score 59;	DB 6;	Length 59;	
Best Local Similarity	100.0%;	Pred. No. 3.3e-23;			
Matches	59;	Conservative 0;	Mismatches 0;	Indels 0;	Gaps 0;
QY	1	CAGCTGAAGAAACGTGTGACACCCCTGCCGAGAAACCGGTGAATCCATCAATAAATG 59			
Db	1	CAGCTGAAGAAACGTGTGACACCCCTGCCGAGAAACCGGTGAATCCATCAATAAATG 59			
RESULT 2	ARI60920/c	59 bp	DNA	linear	PAT 17-OCT-2001
LOCUS	ARI60920	Sequence 11 from patent US 6255281.			
DEFINITION	ARI60920				
ACCESSION	ARI60920.1	GI:16225993			
VERSION					
KEYWORDS					
SOURCE	Unknown.				
ORGANISM	Unknown.				
REFERENCE	1 (bases 1 to 59)				
AUTHORS	Pilon, A.L., Mukherjee, A.B. and Zhang, Z.				

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	misc_feature							
ORIGIN								
	Query Match	32.2%	Score 19,	DB 2,	Length 62318;			
	Best local Similarity	100.0%;	Pred. No. 5.9;					
	Matches 19; Conservative	0;	Mismatches	0;	Indels	0;	Gaps 0;	
Oy	4 CTGAGAAACTGGTTGACA 22							
Dn	9891 CTCGAAGAATCGTTTACA 9909							
RESULT 4								
LOCUS	AC117645	65394 bp	DNA	linear	HNG 30-AUG-2002			
DEFINITION	Mus musculus clone RP23-238F4, LOW-PASS SEQUENCE SAMPLING.							
ACCESSION	AC117645							
VERSION	AC117645.5 GI:22549559							
KEYWORDS	HNG, HNGS PHASE0.							
SOURCE	Mus musculus (house mouse)							
ORGANISM	Mus musculus							
REFERENCE	Eukaryota; Metazoa; Chordata; Craniata; Vertebrata; Euteleostomi; Mammalia; Eutheria; Rodentia; Sciurognathi; Muridae; Murinae; Mus. 1 (bases 1 to 65394)							
AUTHORS	Birren,B., Nusbaum,C. and Lander,E.							
TITLE	Mus musculus, clone RP23-238F4							
JOURNAL	Unpublished							
REFERENCE	2 (bases 1 to 65394)							
	Birren,B., Linton,L., Nusbaum,C., Lander,E., All,A., Allen,N., Anderson,S., Barna,N., Bastien,V., Bloom,T., Boguslavsky,I., Boukhalter,B., Brown,A., Camarata,A., Campopiano,A., Chang,J., Chazarro,B., Choepel,Y., Colangelo,M., Collins,S., Collymore,A., Cook,A., Cooke,P., DeArrellano,K., Dewar,K., Diaz,J.S., Dodge,S., Faró,S., Ferreira,P., Fitzhugh,W., Gage,D., Galagan,J., Gardyna,S., Grinde,S., Gord,S., Goeyette,M., Graham,L., Grand-Pierre,N., Hagos,B., Horton,L., Hulme,W., Iliev,I., Johnson,R., Jones,C., Kamat,A., Karatas,A., Kellis,C., Larocque,K., Lamazares,R., Landers,T., Lehoczeky,J., Levine,R., Lindblad-Toh,K., Liu,G., Maclean,C., MacDonald,P., Major,J., Marquis,N., Matthews,C., McCarthy,M., McEwan,P., McKernan,K., Meldrum,J., Menues,L., Mishra,T., Mlenga,V., Murphy,T., Naylor,J., Nguyen,C., Nicol,R., Norbu,C., Norman,C.H., O'Connor,T., O'Donnell,P., O'Neill,D., Oliver,U., Peterson,K., Phunkiang,P., Pierre,N., Pollara,V., Raymond,C., Retta,R., Rieback,W., Riley,R., Rise,C., Rogov,P., Roman,J., Rossetti,M., Roy,A., Santos,R., Schauer,S., Schniback,R., Seaman,S., Severy,P., Spencer,B., Searge-Thomann,N., Stojanovic,N., Strassman,N., Subramanian,A., Talamas,J., Tesfaye,S., Theodore,J., Topham,K., Travers,M., Travis,N., Trigilio,J., Vassiliev,H., Viel,R., Vo,A., Wilson,B., Wu,X., Wyman,D., Ye,W.J., Young,G., Zainoun,J., Zembek,L., Zimmer,A. and Zody,M.							
TITLE	Direct Submission							
JOURNAL	Submitted (10-FEB-2002) Whitehead Institute/MIT Center for Genome Research, 320 Charles Street, Cambridge, MA 02141, USA							
REFERENCE	3 (bases 1 to 65394)							
AUTHORS	Birren,B., Nusbaum,C., Lander,E., All,A., Allen,N., Anderson,S., Barna,N., Bastien,V., Bloom,T., Boguslavsky,I., Boukhalter,B., Camarata,A., Chang,J., Chazarro,B., Choepel,Y., Collymore,A., Cook,A., Cooke,P., DeArrellano,K., Dewar,K., Diaz,J.S., Dodge,S., Faró,S., Ferreira,P., Fitzgerald,M., Gage,D., Galagan,J., Gardyna,S., Gord,S., Graham,L., Grand-Pierre,N., Hagos,B., Horton,L., Hulme,W., Iliev,I., Johnson,R., Jones,C., Kamat,A., Karatas,A., Kellis,C., Landers,T., Levine,R., Lindblad-Toh,K., Liu,G., Maclean,C., MacDonald,P., Major,J., Marquis,N., Matthews,C., McCarthy,M., McEwan,P., McKernan,K., Meldrum,J., Menues,L., Mishra,T., Mlenga,V., Murphy,T., Naylor,J., Nguyen,C., Nicol,R., Norbu,C., Norman,C.H., O'Connor,T., O'Donnell,P., O'Neill,D., Oliver,U., Peterson,K., Phunkiang,P., Pierre,N., Pollara,V., Raymond,C., Retta,R., Rieback,W., Riley,R., Rise,C., Rogov,P., Roman,J., Rossetti,M., Roy,A., Santos,R., Schauer,S., Schniback,R., Seaman,S., Severy,P., Spencer,B., Searge-Thomann,N., Stojanovic,N., Strassman,N., Subramanian,A., Talamas,J., Tesfaye,S., Theodore,J., Topham,K., Travers,M., Travis,N., Trigilio,J., Vassiliev,H., Viel,R., Vo,A., Wilson,B., Wu,X., Wyman,D., Ye,W.J., Young,G., Zainoun,J., Zembek,L., Zimmer,A. and Zody,M.							

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TITLE
JOURNAL
COMMENT

* McCarty,M., Meldrim,J., Marcus,L., Mhova,T., Mlenga,V.,
* Murphy,T., Naylor,J., Nguyen,C., Nicol,J., Norbu,C., Norman,C.H.,
* O'Connor,T., O'Donnell,P., O'Neill,D., Oliver,J., Peterson,K.,
* Phunkhara,P., Pierre,N., Raymond,C., Retta,R., Rise,C., Rogov,P.,
* Roman,J., Roy,A., Schauer,S., Schnuback,K., Seaman,S., Severy,P.,
* Smith,C., Spencer,B., Stange-Thomas,N., Stojanovic,N., Talamas,J.,
* Testa,E.S., Theodore,D., Topham,K., Travers,M., Vassiliev,H.,
* Viel,R., Vo,A., Wilson,B., Wu,X., Wyman,D., Young,G., Zainoun,J.,
* Zemek,L., Zimmer,A. and Zody,M.
Direct Submission
Submitted (30-Aug-2002) Whitehead Institute/MIT Center for Genome
Research, 320 Charles Street, Cambridge, MA 02141, USA
On Aug 30, 2002 this sequence version replaced gi:122474966.
All repeats were identified using RepeatMasker:
Smit,A.F.A. & Green, P. (1996-1997)
http://ftp.genome.washington.edu/RM/RepeatMasker.html
Genome Center
Center: Whitehead Institute/ MIT Center for Genome Research
Center code: WIBR
Web site: http://www-seq.wi.mit.edu
Contact: sequence_submissions@genome.wi.mit.edu
Project Information
Center project name: L23587
Center clone name: 238_F_4

* NOTE: This record contains 81 individual
* sequencing reads that have not been assembled into
* contigs. Runs of N are used to separate the reads
* and the order in which they appear is completely
* arbitrary. Low-pass sequence sampling is useful for
* identifying clones that may be gene-rich and allows
* overlap relationships among clones to be deduced.
* However, it should not be assumed that this clone
* will be sequenced to completion. In the event that
* the record is updated, the accession number will
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12939 13038: gap of 100 bp
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complete sequence.
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VERSION
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KEYWORDS
Oryza sativa (japonica cultivar-group)
Oryza sativa (japonica cultivar-group)
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Ehrhartoideae; Oryzaceae; Oryza.
REFERENCE
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Feng, Q., Zhang, Y., Hao, P., Wang, S., Fu, G., Huang, Y., Li, Y., Zhu, J.,
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Weng, Q., Zhang, L., Lu, Y., Mu, J., Lu, Y., Zhang, L. S., Yu, Z., Fan, D.,
Liu, X., Lu, T., Li, C., Wu, Y., Sun, T., Lai, H., Li, T., Hu, H., Guan, J.,
Wu, M., Zhang, R., Zhou, B., Chen, Z., Chen, Z., Jin, Z., Wang, R.,
Yin, H., Cai, Z., Ren, S., Lv, G., Gu, W., Zhu, G., Tu, Y., Jia, D.,
Zhang, Y., Chen, T., Kang, H., Chen, X., Shao, C., Sun, Y., Hu, Q.,
Zhang, X., Zhang, W., Wang, L., Ding, C., Sheng, H., Gu, J., Chen, S.,
Ni, L., Zhu, F., Chen, W., Lan, L., Lai, Y., Cheng, Z., Gu, M., Jiang, J.,
Li, O., Hong, G., Xue, Y., and Han, B.
Sequence and analysis of rice chromosome 4
JOURNAL
NATURE 420 (6913), 316-320 (2002)
MEDLINE
2237377
PUBMED
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REFERENCE
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Fu, G., Wang, S. Y., Ren, S. X., Lv, G., Lin, G., Gu, W. Q., Zhu, G. P.,
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Zhu, F. H., Han, B., Feng, Q., Huang, Y. C., Li, Y., Zhu, J. J., Zhao, Q.,
Hu, X., Liu, Y. L., Mu, J., Yu, Z., Chen, L., Fan, D. L., Wang, Q. J.,
Zhang, L., Lu, Y. Q., Yu, S. L., Liu, X. H., Lu, T. T., Zhang, Y. J., Lu, Y.,
Li, C., Li, T., Zhang, Y., Hu, H., Jia, P. X., Qian, Y. M., Ying, K.,
Zhou, B., Chen, Z. H., Hao, P., Zhang, L., Wu, M., Zhang, R. Q., Guan, J. P.,
and Hong, G. F.
Direct Submission
Submitted (08-SEP-2001) Han Bin, National Center for Gene Research,
Chinese Academy of sciences, 5004 Cao Bao Road, Shanghai 200233,
CHINA. E-mail enquiries: bhan@ncgr.ac.cn. Clone requests:
bhan@ncgr.ac.cn
Oryza sativa japonica (nipprobare) genomic DNA, chromosome 4, BAC
clone: OSJNB0067920.
On Jul 8, 2003 this sequence version replaced gi:21740578.
Web site: http://www.ncgr.ac.cn
----- Summary Statistics
Assembly program: phrap
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This is a complete sequence.
Genes were identified by a combination of several methods: Gene
prediction programs including Fgenesh (http://www.softberry.com/),
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(http://genemark.biology.gatech.edu/Genemark/), tRNAscan-SE (Sean
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(ftp://ncbi.nlm.nih.gov/blast/db) and the EST database at NCGR.

FEATURES

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Best Local Similarity 100.0%; Pred. No. 5.8;
Matches 19; Conservative 0; Mismatches 0; Indels 0; Gaps 0;

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Rattus norvegicus clone CH230-22601, WORKING DRAFT SEQUENCE, 3
unorderd pieces.

AC111782
AC111782 G1:25073626
HTG: HTGS PHASE1; HTGS DRAFT; HTGS FULLTOP.
Rattus norvegicus (Norway rat)
Rattus norvegicus
Eukaryota; Metazoa; Chordata; Vertebrata; Euteleostomi;
Mammalia; Eutheria; Rodentia; Sciurognathi; Muridae; Murinae;
Rattus.

REFERENCE
AUTHORS
1 (bases 1 to 232344)
Munzy, D., Martz, Metzger, M., Lee, Abramson, S., Adams, C., Alder, J.,
Allen, C., Allen, H., Albrooks, S., Amin, A., Anguiano, D.,
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Wainscoat, G., and Gibbs, R. A.

TITLE
JOURNAL
Direct Submission
Unpublished

REFERENCE
AUTHORS
TITLE
JOURNAL
2 (bases 1 to 232344)
Morley, K.C.
Direct Submission
Submitted (19-FEB-2002) Human Genome Sequencing Center, Department
of Molecular and Human Genetics, Baylor College of Medicine, One
Baylor Plaza, Houston, TX 77030, USA
3 (bases 1 to 232344)
Human Genome Sequencing Consortium.
Submitted (19-NOV-2002) Human Genome Sequencing Center, Department
of Molecular and Human Genetics, Baylor College of Medicine, One
Baylor Plaza, Houston, TX 77030, USA

COMMENT
On Nov 19, 2002 this sequence version replaced gi:23602292.
The sequence in this assembly is a combination of BAC based reads
and whole genome shotgun sequencing reads assembled using Atlas
(http://www.hgsc.bcm.tmc.edu/projects/rat/). Each contig described
in the feature table below represents a scaffold in the Atlas
assembly (a 'contig-scaffold'). Within each contig-scaffold,
individual sequence contigs are ordered and oriented, and separated
by sized gaps filled with Ns to the estimated size. The sequence
may extend beyond the ends of the clone and there may be sequence
contigs within a contig-scaffold that consist entirely of whole
genome shotgun sequence reads. Both end sequences and whole genome
shotgun sequence only contigs will be indicated in the feature
table.

Center: Baylor College of Medicine

Center code: BCM

Web site: http://www.hgsc.bcm.tmc.edu/

Contact: hgsc-help@bcm.tmc.edu

Project Information

Center project name: GOOCG

Center clone name: CH230-22601

Summary Statistics

Assembly program: Phrap; version 0.990329

Consensus quality: 222543 bases at least Q40

Consensus quality: 224732 bases at least Q30

Consensus quality: 226708 bases at least Q20

Estimated insert size: 225765; sum-of-contigs estimation

Quality coverage: 7x in Q20 bases; sum-of-contigs estimation

NOTE: Estimated insert size may differ from sequence length

(see http://www.hgsc.bcm.tmc.edu/docs/genbank_draft_data.html).

NOTE: This is a 'working draft' sequence. It currently

consists of 3 contigs. The true order of the pieces

is not known and their order in this sequence record is

arbitrary. Gaps between the contigs are represented as

* runs of N, but the exact sizes of the gaps are unknown.

* This record will be updated with the finished sequence

* as soon as it is available and the accession number will

* be preserved.

1 226162: contig of 226162 bp in length

226163 226262: gap of unknown length

226263 230757: contig of 4495 bp in length

230758 230857: gap of unknown length

230858 232344: contig of 1487 bp in length.

Location/Qualifiers

1..232344

/organism="Rattus norvegicus"

/mol_type="genomic DNA"

/db_xref="taxon:10116"

/clone="CH230-22601"

1..1538

/note="wgs_end_extension"

clone_end:"77"

1589..2718

/note="wgs_end_extension"

clone_end:"77"

4907..5729

/note="clone_boundary"

clone_end:"T7"

site:

end_sequence:"B2103653"

FEATURES

source

misc_feature

misc_feature

misc_feature

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misc_feature      222851..224016
                  /note="wgs_contig"
misc_feature      224057..226162
                  /note="wgs_contig"
misc_feature      226263..227570
                  /note="wgs_contig"
misc_feature      227922..230757
                  /note="wgs_contig"

ORIGIN
Query Match      32.2%; Score 19; DB 2; Length 232344;
Best Local Similarity 100.0%; Pred. No. 5.8;
Matches 19; Conservative 0; Mismatches 0; Indels 0; Gaps 0;

QY      4 CTGAAGAACTGCTTGACA 22
        |||||
Db      191901 CTGAAGAACTGCTTGACA 191919

RESULT 7
AC128608/c
LOCUS
DEFINITION
ACCESSION
VERSION
KEYWORDS
SOURCE
ORANISM
REFERENCE
AUTHORS

AC128608      246787 bp. DNA linear HTG 09-OCT-2002
Rattus norvegicus clone CH230-320L21, *** SEQUENCING IN PROGRESS
***, 2 unordered pieces.
AC128608
HTG: HTGS PHASE1; HTGS DRAFT; HTGS_ENRICHED.
Rattus norvegicus (Norway rat)
Rattus norvegicus
Eukaryota; Metazoa; Chordata; Vertebrata; Euteleostomi;
Mammalia; Eutheria; Rodentia; Sciurognathi; Muridae; Murinae;
Rattus.
1 (bases 1 to 246787)
Muzny,D,Marle, Netzer,M, Lee, Abramson,S, Adams,C, Alder,J,
Allen,C, Allen,H, Alshbrook,S, Amin,A, Argulano,D,
Ayala-Bechechi,V, Aoyagi,A, Ayodeji,M, Baca,E, Baden,H,
Baldwin,D, Bandaranaike,D, Barber,M, Barnstead,M, Benahmed,F,
Biswal,R, Blair,J, Blankenburg,K, Blyth,P, Brown,M,
Byrant,N, Buhay,C, Burch,P, Burrell,K, Calderon,E,
Cardenas,V, Carter,K, Cavazos,I, Cessat,H, Center,A,
Chacko,J, Chavez,D, Chen,G, Chen,R, Chen,Y, Chen,Z, Chu,J,
Cleveland,C, Cockrell,R, Cox,C, Coyle,M, Cree,A, D'Souza,L,
Davila,M,L, Davis,C, Davy-Carroll,L, De Anda,C, Dederich,D,
Delgado,O, Denson,S, Deramo,C, Ding,Y, Dinh,H, Divya,K,
Diaper,H, Dugan-Rocha,S, Dunn,A, Durbin,K, Duval,B, Evans,K,
Egan,A, Escotto,M, Eugene,C, Evans,C,A, Falls,T, Fan,G,
Fernandez,S, Finley,M, Flagg,N, Forbes,L, Foster,M, Foster,P,
Fraser,C,M, Gabisi,A, Ganta,R, Garcia,A, Garner,T, Garza,M,
Gharghargis,E, Geer,K, Gill,R, Grady,M, Guerra,M, Guevara,W,
Gunnarsson,P, Haaland,W, Hamill,C, Hamilton,C, Hamilton,K,
Harvey,J, Havlak,P, Hawes,A, Henderson,N, Hernandez,J,
Hernandez,R, Hines,S, Hladun,S,L, Hodgson,A, Hogues,M,
Hollins,B, Howells,S, Hulik,S, Hume,J, Idlebird,D, Jackson,A,
Karpelhy,S, Kelly,S, Kelly,S, Khan,Z, King,J, Kovac,C,
Jackson,L, Jacob,L, Jiang,H, Johnson,B, Johnson,R, Jolivet,A,
Kowis,C, Kraft,C,L, Lepow,H, Levan,J, Lewis,L, Li,Z, Liu,J,
Liu,J, Liu,W, Liu,Y, London,P, Longacre,S, Lopez,Z,
Lorenshew,L, Louisedge,H, Lozano,R,J, Lu,X, Ma,D,
Maheshwari,M, Mahindaratne,M, Mahmoud,M, Malloy,K, Mangum,A,
Mangum,B, Mapua,P, Martin,K, Martin,R, Martinez,E,
McInerney,S, McLeod,M,P, McNeill,T,Z, Meenen,E,
Milesavljic,A, Miner,G, Minja,E, Montemayor,J, Moore,S,
Morgan,M, Morris,K, Morris,S, Mundasa,M, Murphy,M, Nair,T,
Nankervis,C, Neal,D, Newton,N, Nguyen,N, Norris,S,
Nwackelamleh,O, Okunolu,G, Olarunpeseoon,A, Pal,S, Parks,K,
Pasternak,S, Paul,H, Perez,A, Perez,L, Pfannkuch,C,
Plopper,F, Poindexter,A, Popovic,D, Primus,E, Pu,L,
Puzo,M, Quiroz,J, Rachlin,E, Reeves,K, Reiter,M,A, Reigh,R,
Riley,B, Reilly,M, Ren,Y, Reuter,M, Richards,S, Riggs,F,
Rivers,C, Rodkey,T, Rojas,A, Rose,M, Rose,R, Ruiz,S,
Sanders,W, Savary,G, Scherer,S, Scott,G, Slatman,S, Shen,H,
Shetty,J, Shvartsbeyn,A, Sisson,I, Sitter,C,D, Smajs,D,
Sneed,A, Sodergren,E, Song,X,Z, Sorelle,R, Sosa,J,
Steinle,M, Strong,R, Sutton,A, Svarek,A, Tabor,P, Taylor,C,
Taylor,T, Thomas,N, Thomas,S, Tingey,A, Trejos,Z, Umami,K,
Valas,R, Vera,V, Villaseana,D, Waldron,J, Walker,B, Wang,J,
Wang,Q, Wang,S, Warren,J, Warren,R, Wei,X, White,F,
Williams,G, Willson,R, Wiczysk,R, Wooden,H, Worley,K,
Wright,D, Wright,R, Wu,J, Yakub,S, Yen,J, Yoon,L, Yoon,V,
Yu,F, Zhang,J, Zhou,X, Zhou,X, Zhao,S, Dunn,D, von
Niederhausen,A, Weiss,R, Smith,D,R, Holt,R,A, Smith,H,O,
Weinstock,G, and Gibbs,R.A.
Direct Submission
Unpublished
2 (bases 1 to 246787)
Worley,K.C.
Direct Submission
Submitted (19-JUL-2002) Human Genome Sequencing Center, Department
of Molecular and Human Genetics, Baylor College of Medicine, One
Baylor Plaza, Houston, TX 77030, USA
3 (bases 1 to 246787)
Rat Genome Sequencing Consortium.
Direct Submission
Submitted (09-OCT-2002) Human Genome Sequencing Center, Department
of Molecular and Human Genetics, Baylor College of Medicine, One
Baylor Plaza, Houston, TX 77030, USA
On Sep 14, 2002 this sequence version replaced gi:21909412.
The sequence in this assembly is a combination of BAC based reads
and whole genome shotgun sequencing reads assembled using Atlas
(http://www.hgsc.bcm.tmc.edu/projects/rat/). As a result, the
sequence may extend beyond the ends of the clone and there may be
contigs that consist entirely of whole genome shotgun sequence
reads. Both end sequences and whole genome shotgun sequence only
contigs will be indicated in the feature table.
----- Genome Center
Center: Baylor College of Medicine
Center code: BCM
Web site: http://www.hgsc.bcm.tmc.edu/
Contact: hgsc-help@bcm.tmc.edu
----- Project Information
Center project name: GYUN
Center clone name: CH230-320L21
----- Summary Statistics
Assembly program: Phrap; version 0.990329
Consensus quality: 219698 bases at least Q40
Consensus quality: 223003 bases at least Q30
Consensus quality: 224973 bases at least Q20
Estimated insert size: 239001; sum-of-contigs estimation
Quality coverage: 4x in Q20 bases; sum-of-contigs estimation
----- NOTE: Estimated insert size may differ from sequence length
(see http://www.hgsc.bcm.tmc.edu/docs/genbankdraftdata.html).
* NOTE: This is a 'working draft' sequence. It currently
* consists of 2 contigs. The true order of the pieces
* is not known and their order in this sequence record is
* arbitrary. Gaps between the contigs are represented as
* runs of N, but the exact sizes of the gaps are unknown.
* This record will be updated with the finished sequence
* as soon as it is available and the accession number will
* be preserved.
* 1 245086; contig of 245086 bp in length
* 245087 245186; gap of unknown length
* 245187 246787; contig of 1601 bp in length.
Location/Qualifiers
1..246787
/organism="Rattus norvegicus"
/mol_type="genomic DNA"
/db_xref="taxon:10116"
/clone="CH230-320L21"
1..1472
/note="wgs_contig"
54443..56179
/note="wgs_contig"
241917..243271
/note="wgs_contig"
243322..245086

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ORIGIN /note="wgs_contig"

Query Match 32.2%; Score 19; DB 2; Length 246787;
Best Local Similarity 100.0%; Pred. No. 5.8;
Matches 19; Conservative 0; Mismatches 0; Indels 0; Gaps 0;

QY 4 CTGAAGAACTGGTTGACA 22
DB 210071 CTGAAGAACTGGTTGACA 210053

RESULT 8

LOCUS A13577 215 bp DNA linear PAT 31-DEC-1993
DEFINITION IGF-II gene.
ACCESSION A13577
VERSION A13577.1 GI:491682
KEYWORDS
SOURCE
ORGANISM
synthetic construct
artificial sequences.

REFERENCE 1 (bases 1 to 215)
AUTHORS Nygren,P.A., Abrahamson,L. and Uhlen,M.
TITLE A recombinant fusion protein, its use and a recombinant vector
JOURNAL Patent: EP 0333691-A 1 20-SEP-1989;
CEMU BIOTECHNIK AB

FEATURES
source location/Qualifiers
1..215
/organism="synthetic construct"
/mol_type="unassigned DNA"
/db_xref="taxon:32630"

gene
CDS
7..213
/gene="IGF II"
7..213
/codon_start=1
/transl_table=11
/protein_id="CA01122.1"
/db_xref="GI:491683"
/translation="MAVRPSETLCGSELVDTLQFVCGDGRGFYFSRPSRVSRRSGIV
EECCFRSCDIALLETVCATPAKSE"

ORIGIN

Query Match 30.5%; Score 18; DB 6; Length 215;
Best Local Similarity 100.0%; Pred. No. 25;
Matches 18; Conservative 0; Mismatches 0; Indels 0; Gaps 0;

QY 11 AACTGGTTGACACCTGC 28
DB 44 AACTGGTTGACACCTGC 61

RESULT 9
LOCUS A13578 215 bp DNA linear PAT 31-DEC-1993
DEFINITION IGF-II gene.
ACCESSION A13578
VERSION A13578.1 GI:489637
KEYWORDS
SOURCE
ORGANISM
synthetic construct
artificial sequences.
1 (bases 1 to 215)
REFERENCE
AUTHORS Nygren,P.A., Abrahamson,L. and Uhlen,M.
TITLE A recombinant fusion protein, its use and a recombinant vector
JOURNAL Patent: EP 0333691-A 2 20-SEP-1989;
CEMU BIOTECHNIK AB

FEATURES
source location/Qualifiers
1..215
/organism="synthetic construct"
/mol_type="unassigned DNA"
/db_xref="taxon:32630"

ORIGIN

Query Match 30.5%; Score 18; DB 6; Length 215;
Best Local Similarity 100.0%; Pred. No. 25;
Matches 18; Conservative 0; Mismatches 0; Indels 0; Gaps 0;

QY 11 AACTGGTTGACACCTGC 28
DB 176 AACTGGTTGACACCTGC 159

RESULT 10

LOCUS SYNBLIFRG 220 bp DNA linear SYN 27-APR-1993
DEFINITION Artificial bovine insulin-like growth factor 2 gene, complete cds.
ACCESSION M60420
VERSION M60420.1 GI:208029
KEYWORDS insulin-like growth factor II.
SOURCE
ORGANISM
synthetic construct
artificial sequences.
1 (bases 1 to 220)
REFERENCE
AUTHORS Easton,A.M., Gierse,J.K., Seetharam,R., Klein,B.K. and Kottis,C.E.
TITLE Production of bovine insulin-like growth factor 2 (BIGF2) in
JOURNAL Escherichia coli
MEDLINE Gene 101 (2), 291-295 (1991)
PUBMED 91276286
COMMENT Original source text: Synthetic DNA.
FEATURES
source location/Qualifiers
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/organism="synthetic construct"
/mol_type="genomic DNA"
/db_xref="taxon:32630"

CDS
7..216
/note="bovine"
/codon_start=1
/transl_table=11
/product="insulin-like growth factor 2"
/protein_id="AA72971.1"
/db_xref="GI:208030"
/translation="MAVRPSETLCGSELVDTLQFVCGDGRGFYFSRPSRVSRRSGIV
VECCFRSCDIALLETVCATPAKSE"

ORIGIN

Query Match 30.5%; Score 18; DB 12; Length 220;
Best Local Similarity 100.0%; Pred. No. 25;
Matches 18; Conservative 0; Mismatches 0; Indels 0; Gaps 0;

QY 11 AACTGGTTGACACCTGC 28
DB 47 AACTGGTTGACACCTGC 64

RESULT 11
LOCUS AK073198 1234 bp mRNA linear PLN 24-JUL-2003
DEFINITION Oryza sativa (japonica cultivar-group) cDNA clone:J033022115, full
insert sequence.
ACCESSION AK073198
VERSION AK073198.1 GI:32983221
KEYWORDS
SOURCE
ORGANISM
Oryza sativa (japonica cultivar-group)
Oryza sativa (japonica cultivar-group)
Eukaryota; Viridiplantae; Streptophyta; Embryophyta; Tracheophyta;
Spermatophyta; Magnoliophyta; Liliopsida; Poales; Poaceae;
Ehrhartoideae; Oryzaceae; Oryza.

REFERENCE
AUTHORS The Rice Full-Length cDNA Consortium, National Institute of
Agrobiological Sciences Rice Full-Length cDNA Project Team,
Kikuchi,S., Satoh,K., Nagata,T., Kawagashira,N., Doi,K.,
Kishimoto,N., Yatake,U., Ishikawa,M., Yamada,H., Ooka,H., Hotra,T.,
Kojima,K., Namiki,T., Ohneda,E., Yahagi,W., Suzuki,K., Li,C.,


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CDS
/gene="trpg"
/length=5185
/gene="trpg"
/codon_start=1
/transl_table=1
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/protein_id="AA80553.1"
/db_xref="GI:151531"
/translation="MLNMDNDYSPFYNNVYVIGLGAFTKIRNDENTIAQIAALN
PERIVSPGCTSEAGVSIPLAIFGAKPLIGLCHGHSIQAGVVRARQVMA
GKTSFVHRDGVGTGNNPLTVTRYSILVKKETLPDCLEVTAMTAHDSVDELMG
LRHKTINIEGVQFPEISILTEQHELPAFLKQGGRR"
5195..6244
/gene="trpg"
/length=6244
/gene="trpg"
/codon_start=1
/transl_table=1
/product="phosphoribosyl transferase"
/protein_id="AA80554.1"
/db_xref="GI:1052829"
/translation="MDIKSALSRTVGLDITTEPRDYMNRQIMTGOCTENAGPILMG
MKMSSTDEIVGAVSWRRLADKVEIKSLDGVYDIYCGDGNAPNYSSTSPVLA
AACTVAKHGKRAVSGKSGKDLLEAGTYINLPTVACISLIGIRPRASHSA
MKHAAQPRDGLKTLFNMGLPLTPAGVHGVVPAQTLCPLEAVLQRLSKHVL
VHSGKDIDERSLAAPTFVALKNGBITEVWEDEDGMKSGSLHGLAVENPQASLEL
IRDALGRKIDENQKAAEMTVINAGALVADHAMTLAOGVELAHVDVLTGLAMEKLO
ELGAFVAFKVENEA"
6241..7074
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/length=7074
/gene="trpg"
/codon_start=1
/transl_table=1
/product="indoleglycerol phosphate synthetase"
/protein_id="AA80555.1"
/db_xref="GI:1052830"
/translation="MSVPTVLEFRIARFOEVARSAVSLSAELERLAKTADAPRGA
NALIFQAKRKPATVIAETIKKASPSKQITREHFPATVAVSEKGCATCISVLDVYF
QADVILQQAARAVSLVPIRKQVNDVPIQVEARAGADCVLIVSLDDKADAA
TAKDVGDLVAVHDGDELEKLTDTPLVANNRLTFEVSLETTDLPLRIARD
RLATSEGLINRADVELMAINEVYSLVGAFAKAEQPGLELQRFEPPEQKXTQVPL
D"

ORIGIN
Query Match 30.5% Score 18; DB 1; Length 7419;
Best Local Similarity 100.0%; Pred. NO. 23;
Matches 18; Conservative 0; Mismatches 0; Indels 0; Gaps 0;

QY 19 GACACCTGCGCAGAAA 36
|||||
Db 388 GACACCTGCGCAGAAA 405

RESULT 15
AC004625/c 42499 bp DNA linear PLN 11-MAR-2002
LOCUS Arabidopsis thaliana chromosome 2 clone T26013 map C1C11C08,
DEFINITION complete sequence.
ACCESSION AC004625
VERSION AC004625.3 GI:20197200
KEYWORDS HTG.
SOURCE Arabidopsis thaliana (thale cress)
ORGANISM Arabidopsis thaliana
Eukaryota; Viridiplantae; Streptophyta; Embryophyta; Tracheophyta;
Spermatophyta; Magnoliophyta; eudicotyledons; core eudicots;
rosids; eurosids II; Brassicales; Brassicaceae; Arabidopsis.
1 (bases 1 to 42499)
REFERENCE Rounsley,S.D., Lin,X., Ketchum,K.A., Crosby,M.L., Brandon,R.C.,
AUTHORS Sykes,S.M., Kaul,S., Mason,T.M., Kellavage,A.R., Adams,M.D.,
Somerville,C.R. and Venter,J.C.
JOURNAL Unpublished
REFERENCE 2 (bases 1 to 42499)

AUTHORS Lin,X.
TITLE Direct Submission
JOURNAL Submitted (09-MAR-2000) The Institute for Genomic Research, 9712
Medical Center Dr., Rockville, MD 20850, USA
REFERENCE 3 (bases 1 to 42499)
AUTHORS Town,C.D. and Kaul,S.
TITLE Direct Submission
JOURNAL Submitted (27-FEB-2002) The Institute for Genomic Research, 9712
Medical Center Dr., Rockville, MD 20850, USA, cdcm@tigr.org
On Apr 18, 2002 this sequence version replaced gi:6598427.
FEATURES
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1..42499
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/mol_type="Genomic DNA"
/cultivar="Columbia"
/db_xref="taxon:3702"
/chromosome="2"
/map="C1C11C08"
/clone="T26013"
/complement(1..5112)
/overlap with BAC clone F13H10 (AC005662:1..5112)."
2683..3649
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/note="synonym: T26013.2; identical to GB D30719;
supported by full length cDNA; Ceres: 31388"
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/codon_start=1
/product="ERD15 protein"
/protein_id="AAC33728.1"
/db_xref="GI:3241941"
/translation="MAWVSGRSTLNPADLFLPAVRYQVEDFSPEMWOLVTSWYF
DWMYSQQQADGDFGDMGENENGSHIDVADLPLRSFDDMDMDFDPTDAEBOGDF
GMYVQAPSEFGKNGEMVTKSGNRSPIVEPAKYAKEPKAWGNQVAAAPENH
QPR"
4440..10934
/gene="At2g41440"
/note="synonym: T26013.3; predicted by genescan"
join(<4440..4488,4522..4676,4969..5123,5491..5528,
5605..5701,5791..5832,6102..6175,6573..6615,6694..6773,
6869..7010,7136..7268,8763..8916,9258..9300,9378..9418,
9516..9615,9739..9789,10210..10289,10327..10421,
10852..>10934)
/gene="At2g41440"
join(4440..4488,4522..4676,4969..5123,5491..5528,
5605..5701,5791..5832,6102..6175,6573..6615,6694..6773,
6869..7010,7136..7268,8763..8916,9258..9300,9378..9418,
9516..9615,9739..9789,10210..10289,10327..10421,
10852..10934)
/gene="At2g41440"
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/product="unknown protein"
/protein_id="AAC33729.1"
/db_xref="GI:3241942"
/translation="MAFKRASPMOSSISYIENSQENSSCVIRFLQVITRQWVR
ISPVNPDPAKSNFPEPTQSKQWKRKIGFSLPDEGRSDDEEAKRSVEQER
ALQFLRILOOLKREVERLRLLIKRMGKDIYSFELLALQSLNDVSIIVEQKEF
VELEEARSPHOKKEASADAGMRGVLVCKFSKSGIDGQEDPGEALPLQMOKEF
EORSSSLOSFPDKLMLFERNNGRELQWTSASLILHIOILARALIDOLGPRQ
EHRAWORGELSPGVRIRMTMLPLPCNVDFSFQSSSDSIILISGRACRDINFG
PSTRLITSMNGRIRIKMLKGRRLDITKKVAILDYGSGSGIIRYDQRPHEHDH
EAVMENBILKRLILTRATKCDITGTPRLLISLKSHTOTALIYKDTFKIKELK
EDDMMNIRLQGLQGFERRRVAEPMIYGLMILKYDVSTFLFWACFFLLFYXTD
RNNGRGLDGVADRFPPGYRYISDRRLTRIVF"
8133..8176
/rpl_family="AT_rich"
8261..8300
/rpl_family="AT_rich"
complement(8423..8563)
/rpl_family="AT_rich"
complement(8590..8611)

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repeat_region
9707..9729
/rpt_family="(TA)n"
/rpt_family="AT_rich"
11427..18324
/gene="At2g41450"
/note="synonym: T26J13.4; predicted by genescan"
join(<11427..11475,11562..11585,11869..11963,12598..12690,
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14281..14339,14419..14606,14666..14706,15252..15372,
15585..15755,16207..16481,16572..16721,16826..16878,
17126..17234,17326..17460,17549..17593,17684..17871,
17960..18101,18181..18324)
/gene="At2g41450"
join(11427..11475,11562..11585,11869..11963,12598..12690,
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14281..14339,14419..14606,14666..14706,15252..15372,
15585..15755,16207..16481,16572..16721,16826..16878,
17126..17234,17326..17460,17549..17593,17684..17871,
17960..18101,18181..18324)
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/codon_start=1
/product="hypothetical protein"
/protein_id="AAC23730.1"
/db_xref="GI:3241943"
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SVGIRTYCADRESEGFMLKQFGLAEVDKAGKGLHISNIRKALCPGSGTLM
LSHNEFTSNLEISSEWYKCEGSPLSARNNSGTPVGDVLRSEGEVYDCTIS
GIRSPWDSITGKNNNVISDQATTPDSEKSTPKLKSMEASLSLOSRIRANNN
NSRIATDPAASAKOSKQNSQVDITDQSLPTICRNKDVCCMKATGIMERPNG
QTRILMDICENKRAFLTEVIRKGLGVTLDGTSHTVITGKRLNLCITLCSG
AMIVSPWLESEYREGFRANESHLHDDYQLKYDTDLKTVLAKARPSVHLKGDY
ICVGNPIELIKTSALIKSAGNVISGVNKKATPIALSLAHGSRVSCHPHNR
SLKTFQILLFVQVDFPKPIISGVDAEMKIVRYLLQMSAILKILTRIVRSWAKC
FLBDCIIOGNSYSRKFLITFLPLSLACLAPLQMSGVNDSVQWDDVPP
GNAVREKSLRSDSCVSLDQSRQAPVRLDKLHPDKLPRITITSTAGLE
CQLPMFVHKVIGVETFFLVFVGTAASPVSVLETTIRGVNVYRTLELSEOKKSI
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ILKLRKGLIRIYAPMVS"
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/note="synonym: T26J13.5; identical to GB:X76912"
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21363..21410,21494..21574,21652..21811,21885..22006)
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/codon_start=1
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/protein_id="AAC23731.1"
/db_xref="GI:3241944"
/translation="MANNVLOFGLQSAIYAKFLVPLRLVSGSFEVGVGSTRSN
KRLMSNATAFINSRKELKIPGAIDONCHOMGSDPDRDEMGLQDKREIEMTV
QELSTRLKLVGVRGKQKELI STLRLHDSNLPPQKETSRSDSVTIKRISNR
EPTDECTNSEADTEHGRKVKOSTEKULKAKVAKAIKAEOKSIMRTGKQOOSKE
EISTSTSELKTEBITSSPQSEPTVLAHKPKOKMRYNPKTMRPPIPESTKCY
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SGDGLKRLSYRIEEMDRTLNRIKELERKSPVLTGDNCAHEEIDIFNPAGNRSAG
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complement(24160..24183)
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/rpt_family="(TA)n"
complement(25382..25598)
/gene="At2g41470"
/note="synonym: T26J13.6"
complement(join(<25382..25532,26138..26199,26283..26315,

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CDS
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28500..28798,29256..29363,29438..299598))
/gene="At2g41470"
complement(join(25382..25532,26138..26199,26283..26315,
26420..26516,26603..26640,26955..27081,27462..27515,
28500..28798,29256..29363,29438..299598))
/gene="At2g41470"
/codon_start=1
/product="unknown protein"
/protein_id="AAC23732.1"
/db_xref="GI:3241945"
/translation="MLRLAIPLPALCSFLPSSARSFTTKPLPIDSIPKPLEN
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IGNTSQSPITTSVAHINRRFHSYQKSMFESSFNIRKMDKRLIHSIFPI
ELNARFESBVOULRYVQDLREKVERLRLFLIRMTGKDDHAFELQLBSRLQV
SRVTDQGEKNIKLEEDERKEAENSDGTRGVLYPCNRRNNGRLEQMSRDLISL
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29864..29901
/rpt_family="(TA)n"
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29864..29901
/rpt_family="AT_rich"
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30223..32014
/gene="At2g41480"
/note="synonym: T26J13.7"
complement(join(<30223..30644,31172..31334,31436..31615,
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/gene="At2g41480"
complement(join(30223..30644,31172..31334,31436..31615,
31706..32014))
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31706..32014
/gene="At2g41480"
/codon_start=1
/product="putative peroxidase"
/protein_id="AAC23733.1"
/db_xref="GI:3241946"
/translation="MVCYERYKVLLEHGMILNHTAQLVRKQVLYGKYIMIM
LVIVGKVRQLKNGVYISGCKKASIVRSYVESHFSDPISGLRLRHHDFV

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Query Match 30.5%; Score 18; DB 8; Length 42499;
Best Local Similarity 100.0%; Pred. No. 23;
Matches 18; Mismatches 0; Indels 0; Gaps 0;
Qy 30 GCAGAAACCGCGTGAATC 47
Db 1361 GCAGAAACCGCGTGAATC 1344
Search completed: December 22, 2004, 23:36:38
Job time : 841.543 secs

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GenCore version 5.1.6
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CM nucleic - nucleic search, using sw model

Run on: December 22, 2004, 19:14:28 / Search time 208.235 Seconds
(without alignments)
1487.336 Million cell updates/sec

Title: US-09-898-616A-3

Perfect score: 59
Sequence: 1 cagctgaagaactgctgtga.....cgtcattcattcataactg 59

Scoring table: OLIGO_NUC
Gapop 60.0, Gapext 60.0

Searched: 4134886 seqs, 2624710521 residues

Word size: 0

Total number of hits satisfying chosen parameters: 8269772

Minimum DB seq length: 0

Maximum DB seq length: 2000000000

Post-processing: Listing first 45 summaries

Database: N_Geneseq_23Sep04:*

1: geneseqn1980a:*\n2: geneseqn1990a:*\n3: geneseqn2000a:*\n4: geneseqn2001a:*\n5: geneseqn2001b:*\n6: geneseqn2002a:*\n7: geneseqn2002b:*\n8: geneseqn2003a:*\n9: geneseqn2003b:*\n10: geneseqn2003c:*\n11: geneseqn2003d:*\n12: geneseqn2004a:*

Pred. No. is the number of results predicted by chance to have a score greater than or equal to the result being printed, and is derived by analysis of the total score distribution.

SUMMARIES

Result No.	Score	Query Match	Length	ID	Description
1	59	100.0	59	9 ABZ58372	ABZ58372 Human ute
2	59	100.0	59	12 ADL27628	ADL27628 Recombina
3	37	62.7	59	9 ABZ58375	ABZ58375 Human ute
4	37	62.7	59	12 ADL27631	ADL27631 Recombina
5	18	30.5	215	1 AAN90906	AAN90906 Synthetic
6	17	28.8	195	6 ABR76145	ABR76145 Bacillus
7	17	28.8	993	10 ADC24050	ADC24050 DNA seque
8	17	28.8	993	12 ADH36151	ADH36151 Chemical
9	17	28.8	993	12 ADG93852	ADG93852 Nitriase
10	17	28.8	993	12 ADI62449	ADI62449 DNA encod
11	17	28.8	993	12 ADI64570	ADI64570 DNA encod
12	17	28.8	3530	4 ABJ08480	ABJ08480 Drosophi
13	17	28.8	5568	10 ABX08140	ABX08140 S. pneumo
14	17	28.8	5568	12 ADM91810	ADM91810 S. pneumo
15	17	28.8	14736	2 AAV52304	AAV52304 Streptoco
16	17	28.8	62598	10 ABZ56454_21	ABZ56454 Human ute
17	16	27.1	60	9 ABZ58374	ABZ58374 Human ute
18	16	27.1	60	12 ADL27630	ADL27630 Recombina
19	16	27.1	311	4 AAI01419	AAI01419 Human rep
20	16	27.1	377	8 ABX55377	ABX55377 Bovine ES
21	16	27.1	603	11 ABD04549	ABD04549 Pseudomon

ALIGNMENTS

22	16	27.1	639	10	ADB69621	ADB69621 C. neofo
23	16	27.1	659	3	AAFO8058	AAFO8058 Fusarium
24	16	27.1	667	4	ABQ75940	ABQ75940 CHD activ
25	16	27.1	743	10	ADB69260	ADB69260 C. neofo
26	16	27.1	933	11	ABD02206	ABD02206 Pseudomon
27	16	27.1	1346	5	AAST73783	AAST73783 DNA encod
28	16	27.1	1769	6	ABN98233	ABN98233 Arabidops
29	16	27.1	2142	11	ABD02265	ABD02265 Pseudomon
30	16	27.1	2658	5	AAJ90260	AAJ90260 DNA encod
31	16	27.1	2658	5	AAJ94408	AAJ94408 DNA encod
32	16	27.1	2658	5	AAJ73288	AAJ73288 DNA encod
33	16	27.1	2659	5	AAJ88462	AAJ88462 DNA encod
34	16	27.1	2743	10	ADB68899	ADB68899 C. neofo
35	16	27.1	2874	8	ACAI8781	ACAI8781 Prokaryot
36	16	27.1	3056	3	AAJ42758	AAJ42758 Arabidops
37	16	27.1	3108	4	ABL24046	ABL24046 Drosophi
38	16	27.1	3163	10	ADE25034	ADE25034 Plant gro
39	16	27.1	4221	12	ADO57346	ADO57346 DNA encod
40	16	27.1	5603	3	AAJ55314	AAJ55314 Human act
41	16	27.1	6564	3	AAJ55314	AAJ55314 Human act
42	16	27.1	10327	4	ABL12538	ABL12538 Drosophi
43	16	27.1	11204	3	AAJ55339	AAJ55339 Human act
44	16	27.1	11204	6	ABJ73286	ABJ73286 DNA encod
45	16	27.1	21469	4	AAK89568	AAK89568 Human dig

RESULT 1

ABZ58372	ABZ58372 standard; DNA; 59 BP.
AC	ABZ58372;
XX	
DT	28-APR-2003 (first entry)
DE	Human uteroglobin synthetic gene oligonucleotide 3.
XX	
XX	Human; uteroglobin; respiratory distress; antiinflammatory; antifibrotic;
KW	antiinflammatory; antiasthmatic; nephroprotectic; antithromatic;
KX	antiarthritic; ss.
XX	
OS	Homo sapiens.
XX	
OS	Synthetic.
XX	
PN	WO2003003979-A2.
XX	
PD	16-JAN-2003.
XX	
PF	02-JUL-2002; 2002WO-US020836.
XX	
PR	02-JUL-2001; 2001US-00898616.
PA	(CLAR-) CLARAGEN INC.
XX	
PI	Pilon AL, Welch RE;
XX	
DR	WPI, 2003-221527/21.
XX	
PT	Bacterial expression system for producing recombinant human uteroglobin
PT	for treating inflammatory and fibrotic conditions, comprises a synthetic
PT	gene which codes for human uteroglobin.
XX	
PS	Claim 1, Page 33; 127pp; English.
XX	
CC	The present sequence is that of oligonucleotide 3, which was used in the
CC	construction of a synthetic gene for the production of human uteroglobin
CC	(hug) in bacteria. Oligonucleotides 1-4 (see ABZ58370-73) were used to
CC	assemble the coding strand and oligonucleotides 5-8 (see ABZ58374-77) the
CC	complementary strand. The gene was assembled by annealing and ligation of
CC	the oligonucleotides. Because mature native hug has glutamic acid at its
CC	N-terminus, an initiator methionine was added to the N-terminus, and

CC codon usage was optimised for expression in bacteria. In an example from
CC the invention, the synthetic gene was cloned into plasmid pCG12 (see
CC AB258378) and recombinant hUG (see ABP72259) was produced in *Escherichia*
CC coli strain CG12. The invention relates generally to the production of
CC recombinant hUG by bacterial expression, protein purification and scaled-
CC up production according to current good manufacturing practices. The
CC recombinant hUG is useful for the treatment of inflammatory and fibrotic
CC conditions, such as neonatal respiratory distress syndrome and
CC bronchopulmonary dysplasia. It may also be used to treat conditions
CC associated with elevated phospholipase A2 levels such as pancreatitis,
CC acute renal failure, rheumatoid arthritis and asthma
CC
XX
SQ Sequence 59 BP; 19 A; 17 C; 13 G; 10 T; 0 U; 0 Other;
Query Match 100.0%; Score 59; DB 9; Length 59;
Best Local Similarity 100.0%; Pred. No. 7.3e-22;
Matches 59; Conservative 0; Mismatches 0; Indels 0; Gaps 0;
Dy 1 CAGCTGAAGAACTGGTTGACACCTGCGCGAGAAACCGCGTGAATCCATCATTAACCTG 59
Db 1 CAGCTGAAGAACTGGTTGACACCTGCGCGAGAAACCGCGTGAATCCATCATTAACCTG 59
RESULT 2
ADL27628
ID ADL27628 standard; DNA; 59 BP.
XX
AC ADL27628;
XX
DT 20-MAY-2004 (first entry)
XX
DE Recombinant human uteroglobin, rhUG, coding oligonucleotide #3.
XX
XX Human; ss; recombinant human uteroglobin; rhUG;
XX bacterial expression system; rhUG master cell bank;
XX rhUG research seed bank; anti-inflammatory; secretory phospholipase A 2;
XX fibronectin; respiratory distress; inflammation; fibrotic disease.
XX
OS Homo sapiens.
OS Synthetic.
OS
PN US2003207795-A1.
XX
PD 06-NOV-2003.
XX
PE 02-JUL-2002; 2002US-00187498.
XX
PF 28-MAY-1997; 97US-00864357.
XX
PR 02-JUL-2001; 2001US-00898616.
XX
PA (PILON) PILON A L.
XX (WELC) WELCH R W.
XX
PI Pilon AL, Welch RW;
XX
DR WPI; 2004-051527/05.
XX
PT Bacterial expression system for production of recombinant human
PT uteroglobin comprising synthetic gene or human cDNA sequence which codes
PT for human uteroglobin.
XX
XX Claim 1; SEQ ID NO 3; 64bp; English.
XX
XX The invention relates to a bacterial expression system for the production
XX of recombinant human uteroglobin (rhUG), comprising a synthetic gene or
XX human cDNA sequence which codes for human UG, constructed from the
XX oligonucleotides appearing as ADL27626-ADL27629, and which further
XX comprises Met-Ala-Ala at the N-terminus of the sequence. Also included
XX are producing an rhUG master cell bank (comprising inoculating a suitable
XX incubating broth with an aliquot portion of a rhUG research seed bank to
XX form a bacterial culture, incubating the bacterial culture, adding a
XX cryoprotective to the bacterial culture to form a cryopreserved
XX solution, transferring a portion of the cryopreserved solution to a

CC cryovial and storing the cryovial at a temperature below -60 degrees C),
CC expressing rhUG (comprising providing a production seed cell bank culture
CC comprising an expression vector capable of expressing rhUG, inoculating a
CC broth medium with the production seed cell bank culture to form an
CC inoculum, incubating the bacterial culture formed in step (b),
CC inoculating a large scale fermenter with the inoculum formed from the
CC step (c) to form a fermentation culture, incubating the fermentation
CC culture within the large scale fermenter, adding an induction agent to
CC the fermentation culture to induce the expression of rhUG and harvesting
CC the above fermentation culture), purifying rhUG, determining the potency
CC of rhUG in a sample, measuring in vitro anti-inflammatory effect arising
CC from inhibition or blocking of secretory phospholipase A 2 enzymes by
CC rhUG, measuring in vitro binding of rhUG to fibronectin, determining the
CC purity of rhUG, and a pharmaceutical composition comprising a purified
CC rhUG and a carrier or diluent. The bacterial expression system is useful
CC for producing a rhUG research seed bank or a pharmaceutical grade rhUG
CC drug substance. rhUG is safe to administer to a patient in respiratory
CC distress. The rhUG is useful for treating inflammation and fibrotic
CC diseases. The present sequence is a coding strand oligonucleotide used to
CC construct the synthetic rhUG gene.
XX
SQ Sequence 59 BP; 19 A; 17 C; 13 G; 10 T; 0 U; 0 Other;
Query Match 100.0%; Score 59; DB 12; Length 59;
Best Local Similarity 100.0%; Pred. No. 7.3e-22;
Matches 59; Conservative 0; Mismatches 0; Indels 0; Gaps 0;
Dy 1 CAGCTGAAGAACTGGTTGACACCTGCGCGAGAAACCGCGTGAATCCATCATTAACCTG 59
Db 1 CAGCTGAAGAACTGGTTGACACCTGCGCGAGAAACCGCGTGAATCCATCATTAACCTG 59
RESULT 3
ABZ58375/c
ID ABZ58375 standard; DNA; 59 BP.
XX
AC ABZ58375;
XX
DT 28-APR-2003 (first entry)
XX
DE Human uteroglobin synthetic gene oligonucleotide 6.
XX
XX Human; uteroglobin; respiratory distress; antiinflammatory; antifibrotic;
XX antiinflammatory; antiaesthetic; nephrotropic; antirheumatic;
XX antiarthritic; ss.
XX
OS Homo sapiens.
OS Synthetic.
OS
PN WO2003003979-A2.
XX
PD 16-JAN-2003.
XX
PE 02-JUL-2002; 2002WO-US020836.
XX
PR 02-JUL-2001; 2001US-00898616.
XX
PA (CLAR-) CLARAGEN INC.
XX (WELC) WELCH RE;
XX
PI Pilon AL, Welch RE;
XX
DR WPI; 2003-221527/21.
XX
XX Bacterial expression system for producing recombinant human uteroglobin
XX for treating inflammatory and fibrotic conditions, comprises a synthetic
XX gene which codes for human uteroglobin.
XX
XX Example 1; Page 33; 127bp; English.
XX
XX The present sequence is that of oligonucleotide 6, which was used in the
XX construction of a synthetic gene for the production of human uteroglobin
XX (rhUG) in bacteria. Oligonucleotides 1-4 (see ABZ58370-73) were used to
XX assemble the coding strand and oligonucleotides 5-8 (see ABZ58374-77) the

complementary strand. The gene was assembled by annealing and ligation of the oligonucleotides. Because mature rhug has glutamic acid at its N-terminus, an initiator methionine was added to the N-terminus, and codon usage was optimized for expression in bacteria. In an example from the invention, the synthetic gene was cloned into plasmid pCG12 (see AB25378) and recombinant rhug (see ABP72259) was produced in *Escherichia coli* strain CG12. The invention relates generally to the production of recombinant rhug by bacterial expression, protein purification and scaled-up production according to current good manufacturing practices. The recombinant rhug is useful for the treatment of inflammatory and fibrotic conditions, such as neonatal respiratory distress syndrome and bronchopulmonary dysplasia. It may also be used to treat conditions associated with elevated phospholipase A2 levels such as pancreatitis, acute renal failure, rheumatoid arthritis and asthma.

Sequence 59 BP; 9 A; 17 C; 15 G; 18 T; 0 U; 0 Other;

Query Match 62.7%; Score 37; DB 9; Length 59;
Best Local Similarity 100.0%; Pred. No. 4.9e-10;
Matches 37; Conservative 0; Mismatches 0; Indels 0; Gaps 0;

1 CAGCTGAAGAAACTGGTTGACACCCCTGCCGAGAAAC 37
37 CAGCTGAAGAAACTGGTTGACACCCCTGCCGAGAAAC 1

RESULT 4
ADL27631/C
ID ADL27631 standard; DNA; 59 BP.
XX
XX ADL27631:
XX
XX 20-MAY-2004 (first entry)
XX
DE Recombinant human uteroglobin, rhug, non-coding oligonucleotide #2.
XX
XX Human; ss; recombinant human uteroglobin, rhug;
XX bacterial expression system; rhug master cell bank;
XX rhug research seed bank; anti-inflammatory; secretory phospholipase A 2;
XX fibronectin; respiratory distress; inflammation; fibrotic disease.
XX
XX Homo sapiens.
XX Synthetic.
XX
XX US2003207795-A1.
XX
XX 06-NOV-2003.
XX
XX 02-JUL-2002; 2002US-00187498.
XX
XX 28-MAY-1997; 97US-00864357.
XX
XX 02-JUL-2001; 2001US-00898616.
XX
XX (PILLO/) PILLO A L.
XX (WELC/) WELCH R W.
XX
XX Pilon AL, Welch RW;
XX
XX WPI; 2004-051527/05.
XX
XX Bacterial expression system for production of recombinant human
XX uteroglobin comprising synthetic gene or human cDNA sequence which codes
XX for human uteroglobin.
XX
XX Example 1; SEQ ID NO 6; 64bp; English.
XX
XX The invention relates to a bacterial expression system for the production
XX of recombinant human uteroglobin (rhug), comprising a synthetic gene or
XX human cDNA sequence which codes for human uteroglobin, constructed from the
XX oligonucleotides appearing as ADL27626-ADL27629, and which further
XX comprises Met-Ala-Ala at the N-terminus of the sequence. Also included
XX are producing an rhug master cell bank (comprising inoculating a suitable
XX incubating broth with an aliquot portion of a rhug research seed bank to

form a bacterial culture, incubating the bacterial culture, adding a cryoprotective to the bacterial culture to form a cryopreserved solution, transferring a portion of the cryopreserved solution to a cryovial and storing the cryovial at a temperature below -60 degrees C).
expressing rhug (comprising providing a production seed cell bank culture comprising an expression vector capable of expressing rhug; inoculating a broth medium with the production seed cell bank culture to form an inoculum, incubating the bacterial culture formed in step (b),
inoculating a large scale fermenter with the inoculum formed from the step (c) to form a fermentation culture, incubating the fermentation culture within the large scale fermenter, adding an induction agent to the fermentation culture to induce the expression of rhug and harvesting the above fermentation culture), purifying rhug, determining the potency of rhug in a sample, measuring in vitro anti-inflammatory effect arising from inhibition or blocking of secretory phospholipase A 2 enzymes by rhug, measuring in vitro binding of rhug to fibronectin, determining the purity of rhug, and a pharmaceutical composition comprising a purified rhug and a carrier or diluent. The bacterial expression system is useful for producing a rhug research seed bank or a pharmaceutical grade rhug drug substance. rhug is safe to administer to a patient in respiratory distress. The rhug is useful for treating inflammation and fibrotic diseases. The present sequence is a non-coding strand oligonucleotide used to construct the synthetic rhug gene.

Sequence 59 BP; 9 A; 17 C; 15 G; 18 T; 0 U; 0 Other;

Query Match 62.7%; Score 37; DB 12; Length 59;
Best Local Similarity 100.0%; Pred. No. 4.9e-10;
Matches 37; Conservative 0; Mismatches 0; Indels 0; Gaps 0;

1 CAGCTGAAGAAACTGGTTGACACCCCTGCCGAGAAAC 37
37 CAGCTGAAGAAACTGGTTGACACCCCTGCCGAGAAAC 1

RESULT 5
AAN90906
ID AAN90906 standard; DNA; 215 BP.
XX
XX AAN90906:
XX
XX 25-MAR-2003 (revised)
XX 26-JAN-1990 (first entry)
XX
XX Synthetic human IGF-II.
XX
XX Synthetic IGF-II; fusion protein.
XX
XX Homo sapiens.
XX
XX EP333691-A.
XX
XX 20-SEP-1989.
XX
XX 16-MAR-1989; 89EP-00850091.
XX
XX 17-MAR-1988; 88SE-00000981.
XX
XX (CEMU-) CEMU BIOTECHNIK AB.
XX (SEMB-) SEM BIOTECHNIK AB.
XX
XX Nygren PA, Abrahamson L, Uhlen M;
XX
XX WPI; 1989-272436/38.
XX
XX P-PSDB; AAP91389.
XX
XX New recombinant fusion protein comprising desired protein - flanked by
XX IGG-binding domain of staphylococcal protein A and albumin-binding domain
XX of streptococcal protein G.
XX
XX Disclosure, Fig 1; 10pp; English.
XX
XX Synthetic IGF-II was constructed from 22 oligonucleotides. The gene was

designed with EcoRI and HindIII 5' and 3' cohesive ends resp. to facilitate cloning. An N-terminal methionine codon was included to allow cleavage of fusion proteins with CMV to generate native IGF-II. A double TAA stop codon was incorporated at the 3' end, the gene was used to construct a fusion vector with DNA coding for an IgG-binding domain from staphylococcal protein A and an albumin-binding part of streptococcal protein G. (Updated on 25-MAR-2003 to correct PA field.)

Sequence 215 BP, 35 A, 55 C, 53 G, 72 T, 0 U, 0 Other;

Query Match 30.5%; Score 18; DB 1; Length 215;
Best Local Similarity 100.0%; Pred. No. 8.4;
Matches 18; Conservative 0; Mismatches 0; Indels 0; Gaps 0;

11 AACTGTTGACACCCCTGC 28
DB 44 AACTGTTGACACCCCTGC 61

RESULT 6
ABK76145/c
ID ABK76145 standard; DNA, 195 BP.

ABK76145;
13-AUG-2002 (first entry)

Bacillus licheniformis genomic sequence tag (GST) #3436.

Differential gene expression; genomic sequenced tag; GST;
altered culture condition; environmental stress;
physiological provocation; de.

Bacillus licheniformis.

WO200229113-A2.

11-APR-2002.

05-OCT-2001; 2001WO-US01437.

06-OCT-2000; 2000US-00680598.

27-MAR-2001; 2001US-0279526P.

(NOVO) NOVOSYNES BIOTECH INC.

(NOVO) NOVOSYNES AS.

Berka R, Clausen IG;

WPI; 2002-416684/44.

Monitoring differential expression of several genes in first Bacillus cell relative to expression of same genes in one or more second Bacillus cells, by using substrate containing Bacillus genomic sequenced tag array.

Claim 4; SEQ ID NO 3436; 200pp; English.

The invention describes a method of monitoring differential expression of genes in a first Bacillus cell relative to expression of the genes in other Bacillus cells, comprising hybridising labelled nucleic acid probes isolated from Bacillus cells to a substrate containing array of Bacillus genomic sequenced tags (GST), examining the array, and determining relative gene expression by an observed hybridisation reporter signal of a spot in the array. The method is useful for measuring the expression of genes in a first Bacillus cell relative to expression of the same genes in one or more second Bacillus cells. The method is useful for monitoring global expression of several genes from a Bacillus cell, discovering new genes, identifying possible functions of unknown open reading frames and monitoring gene copy number variation and stability. Monitoring changes in expression of genes may be used to provide a representation of the way in which Bacillus cells adapt to changes in culture conditions, environmental stress or other physiological provocation. Extensive follow

-up characterisation is unnecessary, when one spot on an array equals one gene or one open reading frame, since sequence information is available. This sequence represents a genomic sequence tag (GST) used in the method of the invention. Note: The sequence data for this patent did not form part of the printed specification, but was obtained in electronic format directly from WIPO at http://wipo.int/pub/published_pct_sequences

Sequence 195 BP, 34 A, 48 C, 56 G, 57 T, 0 U, 0 Other;

Query Match 28.8%; Score 17; DB 6; Length 195;
Best Local Similarity 100.0%; Pred. No. 29;
Matches 17; Conservative 0; Mismatches 0; Indels 0; Gaps 0;

21 CACCTGCGCGCAGAAGC 37
DB 102 CACCTGCGCGCAGAAGC 86

RESULT 7
ADC24050/c
ID ADC24050 standard; DNA, 993 BP.

ADC24050;

16-DEC-2003 (first entry)

DNA sequence (SegID 317) encoding a nitrilase enzyme.

gene; de; nitrilase; nitrile; cyanohydrin; ammonia; biocatalyst;
enantiotmer; chiral medicine.

Unidentified.

WO2003000840-A2.

03-JAN-2003.

15-MAY-2002; 2002WO-US015983.

21-JUN-2001; 2001US-0300189P.

30-JUN-2001; 2001US-0309006P.

22-JAN-2002; 2002US-0351336P.

(DIVE-) DIVERSA CORP.

(MADP/) MADDEN D.

Madden M, Desantis G, Chaplin JA, Weiner DP, Milan A, Chl E;

Short JM, Burk M;

WPI; 2003-201417/19.

P-PDB; ADC24051.

Novel nitrilase polypeptide, useful for making (R)- or (S)-ethyl-4-cyano-3-hydroxybutyric acid or (R)- or (S)-mandelic acid or (S)- or (R)-phenyl lactic acid derivative and for producing pharmaceutical composition, and food additive.

Claim 1; SEQ ID NO 317; 560pp; English.

This invention relates to nitrilases and the nucleic acids that encode these enzymes thereof. Specifically, it refers to polypeptides that exhibit nitrilase activity, i.e. the ability to directly hydrolyse nitriles or cyanohydrins into their corresponding carboxylic acids and ammonia. Nitrilases have commercial utility as biocatalysts for use in the synthesis of enantiomerically pure aromatic and aliphatic amino acids, as well as hydroxy acids, which are important for the development of chiral medicines. Furthermore, the present invention describes nitrilases, isolated from mesophilic microorganisms, that have improved activity and stability at increased pH and temperature. They are also inexpensive, efficient catalysts, have broad substrate specificity and are capable of chiral differentiation. This polynucleotide is a DNA sequence that encodes a nitrilase enzyme of the invention.

SQ Sequence 993 BP, 188 A, 330 C, 305 G, 170 T, 0 U, 0 Other;

Query Match 28.8%; Score 17; DB 10; Length 993;

Best Local Similarity 100.0%; Pred. No. 30;

Matches 17; Conservative 0; Mismatches 0; Indels 0; Gaps 0;

QY 37 CCGCGTGAATCCATCAT 53
DB 860 CCGCGTGAATCCATCAT 844

RESULT 8
ADH36151/c
ID ADH36151 standard; DNA, 993 BP.

AC ADH36151;

DT 11-MAR-2004 (first entry)

DE Chemical process monitoring-related nitrilase gene sequence SeqID317.

KW Chemical process monitoring; biochemical process monitoring; cyanide;

KW high throughput system; gene; ds.

OS Unidentified.

PN WO2003098187-A2.

PD 27-NOV-2003.

PF 15-MAY-2003; 2003WO-US015639.

PR 15-MAY-2002; 2002US-0380737P.

PA (DIVE-) DIVERSA CORP.

PI Weiner D, Chaplin JA, Chi E, Milan A, Desantis G, Burk MJ;

PT McQuaid J, Siege J;

DR WPI; 2004-142708/14.

DR P-PSDB; ADH36152.

PT Monitoring a chemical or biochemical process comprising providing a

PT reactant comprising a cyanide or a material that can be converted to

PT cyanide or a reactant that generates a cyanide or a material that can be

PT converted to cyanide.

PS Claim 74; SEQ ID NO 317; 277bp; English.

CC This invention relates to a novel method of monitoring chemical or

CC biochemical processes. The method involves providing a reactant

CC comprising cyanide (or a material that can be converted to a cyanide)

CC that generates as a reaction product cyanide or a material that can be

CC converted to cyanide and measuring the concentration of produced cyanide.

CC The method is useful for monitoring a chemical or biochemical process.

CC The method is effective for high throughput systems and is sufficiently

CC sensitive to detect a small amount of product. The present sequence is

CC that of a gene which encodes a nitrilase enzyme which can be used in the

ID ADG93852 standard; DNA, 993 BP.

AC ADG93852;

DT 11-MAR-2004 (first entry)

DE Nitrilase enzyme gene sequence SeqID317.

KW nitrilase; nitrile; carboxylic acid; chemical process; pH; temperature;

KW enantioselective transformation; gene; ds.

OS Unidentified.

PN WO2003097810-A2.

PD 27-NOV-2003.

PF 15-MAY-2003; 2003WO-US015712.

PR 15-MAY-2002; 2002US-00146772.

PR 09-SEP-2002; 2002US-00241742.

PA (DIVE-) DIVERSA CORP.

PI Desantis G, Short JM, Burk M, Wong K, Farwell R, Chatman K;

PT WPI; 2004-090637/09.

DR P-PSDB; ADG93853.

PT New isolated or recombinant nucleic acid encoding a polypeptide having

PT nitrilase activity, useful for screening enantioselective transformation.

PS Claim 44; SEQ ID NO 317; 295bp; English.

CC This invention is related to a novel isolated or recombinant nucleic acid

CC encoding a protein having nitrilase activity. Nitrilases are capable of

CC converting nitriles directly to carboxylic acids and have great

CC potential for use in industrial chemical processes. The isolated

CC nitrilase proteins of the invention have increased activity and stability

CC at increased pH and temperature when compared to those conventionally

CC used. In addition, the nucleic acid of the invention is useful for

CC screening enantioselective transformation. The present sequence is that

CC of a DNA sequence which encodes a nitrilase enzyme of the invention.

SQ Sequence 993 BP, 188 A, 330 C, 305 G, 170 T, 0 U, 0 Other;

Query Match 28.8%; Score 17; DB 12; Length 993;

Best Local Similarity 100.0%; Pred. No. 30;

Matches 17; Conservative 0; Mismatches 0; Indels 0; Gaps 0;

QY 37 CCGCGTGAATCCATCAT 53

DB 860 CCGCGTGAATCCATCAT 844

RESULT 10

AD162449/c

ID AD162449 standard; DNA, 993 BP.

AC AD162449;

DT 22-APR-2004 (first entry)

DE DNA encoding nitrilase polypeptide #159.

KW Atorvastatin; (R)-ethyl 4-cyano-3-hydroxybutyrate;

KW (R)-ethyl 4-cyano-3-hydroxybutyric acid; epichlorohydrin;

KW 3-hydroxyglutaronitrile; 3-hydroxyglutaronitrile;

KW 4-cyano-3-hydroxybutyric acid; ethyl 4-cyano-3-hydroxybutyric acid;

KW mixed hyperlipidaemia; homozygous familial hypercholesterolaemia;

KW antihypertensive; gene; ds.

OS Unidentified.

XX XX
 EN W02003106415-A2.
 PD XX
 PD 24-DEC-2003.
 XX XX
 PF 13-JUN-2003; 2003MO-US018840.
 XX XX
 PR 13-JUN-2002; 2002US-0389317P.
 PR 26-JUN-2002; 2002US-0392994P.
 XX XX
 PA (DIVE-) DIVERSA CORP.
 PI Burk M, Desantis G, Morgan B, Zhu Z;
 XX
 DR MPI; 2004-090821/09.
 DR P-PSDB; ADI62450.
 XX XX
 PT Preparation of atorvastatin comprises catalytic conversion of 3-
 PT hydroxyglutaronitrile by polypeptide with nitrilase activity, converting
 PT obtained 4-cyano-3-hydroxybutyric acid to ethyl-4-cyano-3-hydroxybutyric
 PT acid and forming atorvastatin.
 XX XX
 PS Claim 46; SEQ ID NO 317; 253pp; English.
 XX XX
 CC The present invention relates to a method for preparing an atorvastatin
 CC intermediate known as (R)-ethyl 4-cyano-3-hydroxybutyrate ((R)-ethyl 4-
 CC cyano-3-hydroxybutyric acid). The method comprises optionally converting
 CC epichlorohydrin or equivalent to 3-hydroxyglutaronitrile, catalytic
 CC conversion of 3-hydroxyglutaronitrile or equivalent to 4-cyano-3-
 CC hydroxybutyric acid with a polypeptide having nitrilase activity,
 CC converting 4-cyano-3-hydroxybutyric acid to ethyl-4-cyano-3-hydroxybutyric
 CC acid, and converting this to (R)-ethyl 4-cyano-3-hydroxybutyrate. The
 CC method involves whole cell processes, cell lysate process, "one pot"
 CC processes, and "multi-pot" processes using a variety of parameters.
 CC Atorvastatin is used, in conjunction with dietary restriction, in the
 CC management of hyperlipidemia, including hypercholesterolaemia, mixed
 CC dyslipidemia and homozygous familial hypercholesterolaemia. The present
 CC sequence encodes a nitrilase polypeptide obtained from an environmental
 CC sample.
 XX XX
 SQ Sequence 993 BP; 188 A; 330 C; 305 G; 170 T; 0 U; 0 Other;
 Query Match 28.8%; Score 17; DB 12; Length 993;
 Best Local Similarity 100.0%; Pred. No. 30;
 Matches 17; Conservative 0; Mismatches 0; Indels 0; Gaps 0;
 OY 37 CCGCGTGAATCCATCAT 53
 Db 860 CCGCGTGAATCCATCAT 844
 RESULT 11
 ADI64570/C
 ID ADI64570 standard; DNA; 993 BP.
 XX
 AC ADI64570;
 XX
 DT 22-APR-2004 (first entry)
 XX
 DE DNA encoding nitrilase seq id 159.
 XX
 KW (R)-ethyl 4-cyano-3-hydroxybutyric acid; nitrile hydrolysis;
 KW carboxylic acid; cyanohydrin moiety hydrolysis;
 KW aminonitrile moiety hydrolysis; chiral alpha-hydroxy acid molecule;
 KW chiral amino acid molecule; (S)-ethyl 4-cyano-3-hydroxybutyric acid;
 KW (R)-mandelic acid; (S)-mandelic acid; (S)-phenyl lactic acid derivative;
 KW (R)-phenyl lactic acid derivative; & enantiomeric excess;
 KW & diastereomeric excess; food additive; drug intermediate; ds; nitrilase;
 KW gene.
 XX
 XX Undeidentified.
 OS
 XX
 PN US2004014195-A1.

XX XX
 PD 22-JUN-2004.
 PD XX
 PF 15-MAY-2003; 2003US-00440523.
 XX XX
 PR 29-DEC-1999; 99US-0173609P.
 PR 07-DEC-2000; 2000US-0254414P.
 PR 28-DEC-2000; 2000US-0075129P.
 PR 21-JUN-2001; 2001US-0300189P.
 PR 30-JUL-2001; 2001US-0309006P.
 PR 22-JAN-2002; 2002US-0351336P.
 PR 15-MAY-2002; 2002US-00146772.
 PR 09-SEP-2002; 2002US-00241742.
 XX XX
 PA (DIVE-) DIVERSA CORP.
 PI Desantis G, Short JM, Burk MJ, Wong K, Farwell R, Chatman K;
 XX
 DR MPI; 2004-121569/12.
 DR P-PSDB; ADI64571.
 XX XX
 PT Novel isolated or recombinant polypeptide having nitrilase activity,
 PT useful in production of food additives.
 XX XX
 PS Claim 1; SEQ ID NO 317; 105pp; English.
 XX XX
 CC The invention describes an isolated or recombinant polypeptide (I)
 CC comprising amino acids having a sequence at least 50 % identical to a
 CC sequence (S1) available in electronic form (EC) from the following web
 CC site ftp.segdata.uspo.gov/sequence.html?DocID=2004014195, or its
 CC variants, having one or more mutations at residue 55 Lys, Gly or Glu, at
 CC residue 60 glutamic acid, at residue 111 Ser, their combinations or
 CC fragments. (I) is useful for: producing an (R)-ethyl 4-cyano-3-
 CC hydroxybutyric acid; hydrolysing a nitrile to a carboxylic acid;
 CC hydrolysing cyanohydrin moiety or an aminonitrile moiety; producing a
 CC chiral alpha-hydroxy acid molecule or a chiral amino acid molecule;
 CC producing an (S)-ethyl 4-cyano-3-hydroxybutyric acid; producing an (R)-
 CC mandelic acid or (S)-mandelic acid; producing (S)-phenyl lactic acid
 CC derivative or an (R)-phenyl lactic acid derivative; modifying a molecule;
 CC and for identifying a modified compound. The inventive method is useful
 CC for monitoring or determining & enantiomeric excess or & diastereomeric
 CC excess. (I) is useful in the production of food additives and drug
 CC intermediates. This sequence encodes a nitrilase of the invention.
 XX XX
 SQ Sequence 993 BP; 188 A; 330 C; 305 G; 170 T; 0 U; 0 Other;
 Query Match 28.8%; Score 17; DB 12; Length 993;
 Best Local Similarity 100.0%; Pred. No. 30;
 Matches 17; Conservative 0; Mismatches 0; Indels 0; Gaps 0;
 OY 37 CCGCGTGAATCCATCAT 53
 Db 860 CCGCGTGAATCCATCAT 844
 RESULT 12
 ABL08480
 ID ABL08480 standard; cDNA; 3530 BP.
 XX
 AC ABL08480;
 XX
 DT 26-MAR-2002 (first entry)
 XX
 DE Drosophila melanogaster expressed polynucleotide SEQ ID NO 19922.
 XX
 KW Drosophila; developmental biology; cell signalling; insecticide;
 KW pharmaceutical; gene; ss.
 XX
 OS Drosophila melanogaster.
 XX
 XX
 PN W0200171042-A2.
 XX
 PD 27-SEP-2001.

XX 23-MAR-2001; 2001WO-US009231.
PF 23-MAR-2000; 2000US-0191637P.
PR 11-JUL-2000; 2000US-00614150.
XX (PEKE) PE CORP NY.
XX Venter JC, Adams M, Li PWD, Myers EW;
PI MPI: 2001-655686/75.
DR P-PSDB: ABB64377.
XX New isolated nucleic acid detection reagent for detecting 1000 or more
PT genes from *Drosophila* and for elucidating cell signaling and cell-cell
PR interactions.
XX Claim 1; SEQ ID NO 19922; 21pp + Sequence Listing; English.
XX The invention relates to an isolated nucleic acid detection reagent
CC capable of detecting 1000 or more genes from *Drosophila*. The invention is
CC useful in developmental biology and in elucidating cell signaling and
CC cell-cell interactions in higher eukaryotes for the development of
CC insecticides, therapeutics and pharmaceutical drugs. The invention
CC discloses genomic DNA sequences (AB16176-AB130511), expressed DNA
CC sequences (AB01840-AB16175) and the encoded proteins (AB57737-
CC AB372072). The sequence data for this patent did not form part of the
CC printed specification, but was obtained in electronic format directly
CC from WIPO at ftp.wipo.int/pub/published_pct_sequences
SQ Sequence 3530 BP; 1121 A; 634 C; 690 G; 1085 T; 0 U; 0 Other;
OY Query Match 28.8%; Score 17; DB 4; Length 3530;
Best Local Similarity 100.0%; Pred. No. 32;
Matches 17; Conservative 0; Mismatches 0; Indels 0; Gaps 0;
DB 1786 CTGAGAACTGGTTGA 20
4 CTGAGAACTGGTTGA 20
CTGAGAACTGGTTGA 1802

XX New proteins and nucleic acid molecules from *Streptococcus pneumoniae*,
PT useful as medicaments for treating or preventing a disease or infection
PT due to *Streptococcus bacteria*, such as pneumonia, sepsis, otitis media or
PT ear infection.
XX Claim 6; SEQ ID NO 4855; 56pp; English.
XX The invention relates to a protein comprising or having at least 50%
CC identity to any of the 2469 amino acid sequences, identified in the
CC specification (available on a computer readable format), or its fragment,
CC expressed from 2469 of 2469 identified DNA coding regions from the
CC *Streptococcus pneumoniae* type 4 strain genomic sequence appearing as
CC AB556454. Also included are an antibody which binds one of the proteins,
CC treating a patient by administering the protein, DNA or antibody (in a
CC composition), a kit comprising first and second primers, which are the
CC nucleic acid cited above or fragments between nucleotides 8-100 of a
CC sequence not defined in the specification, for amplifying a target
CC sequence contained within a *Streptococcus pneumoniae* nucleic acid sequence, where
CC the first primer is substantially complementary to the target sequence
CC and the second primer is substantially complementary to the complement of
CC the target sequence, and where the parts of the primers having
CC substantial complementarity define the terminal of the target sequence to
CC be amplified, assay comprising contacting a test compound with the
CC protein, and determining whether the test compound binds to the protein
CC and a *Streptococcus pneumoniae* bacterium, where one or more genes
CC encoding the proteins has been rendered inactive. The proteins, nucleic
CC acid molecules, antibody and compositions are useful as medicaments for
CC treating or preventing a disease or infection due to *Streptococcus*
CC bacteria, particularly *S. pneumoniae*, such as pneumonia, sepsis, otitis
CC media or ear infection. They are also useful in developing vaccines,
CC diagnostics and antibiotics. The methods are useful for identifying
CC immunodominant proteins. The present sequence is one of the 2469
CC identified coding regions from the genomic sequence. Note: The sequence
CC data for this patent did not form part of the printed specification, but
CC was obtained in electronic format directly from WIPO at
CC ftp.wipo.int/pub/published_pct_sequences. (Updated on 27-OCT-2003 to
CC standardise OS field)
SQ Sequence 5568 BP; 2066 A; 873 C; 1182 G; 1447 T; 0 U; 0 Other;
OY Query Match 28.8%; Score 17; DB 10; Length 5568;
Best Local Similarity 100.0%; Pred. No. 32;
Matches 17; Conservative 0; Mismatches 0; Indels 0; Gaps 0;
DB 4309 TGAATCATCTAAACT 58
42 TGAATCATCTAAACT 58
TGAATCATCTAAACT 4293

PA (TUFT) UNIV TUFTS.
 XX
 PI Camilla A, Hava DL,
 XX WPI; 2004-239189/22.
 DR P-PSDB; ADM92047.
 XX
 PT New Streptococcus pneumoniae nucleic acid molecules, useful for
 PT diagnosing, treating and preventing active infections of Streptococcus
 PT pneumoniae.
 XX
 PS Claim 1; SEQ ID NO 7; 123bp; English.
 XX
 CC This invention relates to novel isolated Streptococcus pneumoniae nucleic
 CC acid molecules and the antigenic polypeptides encoded by them. The
 CC invention may be useful for the production of compounds with an
 CC antibacterial activity or for gene therapy. The nucleic acid molecules,
 CC compositions and methods disclosed are useful for treating Streptococcus
 CC pneumoniae infection. The present sequence is that of an S pneumoniae
 CC gene of the invention.
 XX
 SQ Sequence 5568 BP; 2066 A; 873 C; 1182 G; 1447 T; 0 U; 0 Other;
 XX
 Query Match 28.8%; Score 17; DB 12; Length 5568;
 Best Local Similarity 100.0%; Pred. No. 32;
 Matches 17; Conservative 0; Mismatches 0; Indels 0; Gaps 0;
 QY 42 TGAATCCATCATTAACCT 58
 DB 4309 TGAATCCATCATTAACCT 4293
 XX
 RESULT 15
 ID AAV52304/c
 CC AAV52304 standard; DNA; 14736 BP.
 XX
 AC AAV52304;
 XX
 DT 23-OCT-1998 (first entry)
 XX
 DE Streptococcus pneumoniae genome fragment SEQ ID NO:171.
 XX
 KM Streptococcus pneumoniae; S. pneumoniae; genome; diagnosis; assay;
 KM computer readable medium; vaccine; pharmaceutical composition; ds.
 XX
 OS Streptococcus pneumoniae.
 XX
 PN MO9818931-A2.
 XX
 PD 07-MAY-1998.
 XX
 PP 30-OCT-1997; 97MO-US019588.
 XX
 PR 31-OCT-1996; 96US-0029960P.
 XX
 PA (HUMA-) HUMAN GENOME SCI INC.
 XX
 PI Kunsch CA, Choi GH, Dillon PJ, Rosen CA, Barash SC, Fannon M;
 PI Dougherty BA;
 XX
 DR WPI; 1998-272225/24.
 XX
 PT Computer-readable medium with recorded Streptococcus pneumoniae
 PT polynucleotide sequences - useful in diagnostic kits and assays, and
 PT pharmaceutical compositions and vaccines for Streptococcus pneumoniae.
 XX
 PS Claim 1; Page 1085-1094; 1409bp; English.
 XX
 CC The present invention describes a computer readable medium which has the
 CC nucleotide sequences SEQ ID NO:1 to 391 (AAV52134 to AAV52524) recorded
 CC on it, or a representative fragment or a sequence at least 95% identical
 CC to SEQ ID NO:1 to 391. The nucleotide sequences depicted in SEQ ID NO:1
 CC to 391 (AAV52134 to AAV52524) are genomic fragments from Streptococcus

CC pneumoniae. The present invention also describes an isolated nucleic acid
 CC molecule encoding a homologue of any of the fragments of the S. pneumoniae
 CC genome (SEQ ID NO:1 to 391) where the nucleic acid molecule is produced
 CC by a process comprising: (a) screening a genomic DNA library using as a
 CC probe a target sequence defined by any of the sequences in SEQ ID NO:1 to
 CC 391, identifying members of the library which contain sequences that
 CC hybridize to the target sequence and isolating the nucleic acid molecules
 CC from the members; or (b) isolating mRNA, DNA or cDNA produced from an
 CC organism, amplifying nucleic acid molecules whose nucleotide sequence is
 CC homologous to amplification primers derived from the fragment of the S.
 CC pneumoniae genome to prime the amplification and isolating the amplified
 CC sequences. The computer readable medium can be used in a computer-based
 CC system for identifying fragments of the S. pneumoniae genome of
 CC commercial importance, or expression modulating fragments of the S.
 CC pneumoniae genome. Products from the present invention can be used in
 CC diagnosis kits and assays, and pharmaceutical compositions and vaccines
 CC for S. pneumoniae
 XX
 SQ Sequence 14736 BP; 4665 A; 2568 C; 3255 G; 4247 T; 0 U; 1 Other;
 XX
 Query Match 28.8%; Score 17; DB 2; Length 14736;
 Best Local Similarity 100.0%; Pred. No. 33;
 Matches 17; Conservative 0; Mismatches 0; Indels 0; Gaps 0;
 QY 42 TGAATCCATCATTAACCT 58
 DB 13144 TGAATCCATCATTAACCT 13128
 XX
 Search completed: December 22, 2004, 22:44:17
 Job time : 213.485 secs

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OM nucleic - nucleic search, using sw model

Run on: December 22, 2004, 22:08:48 / Search time 47.5204 Seconds
(without alignments)
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Title: US-09-898-616A-3

Perfect score: 59

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Gapop 60.0, Gapext 60.0

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Total number of hits satisfying chosen parameters: 1649014

Minimum DB seq length: 0
Maximum DB seq length: 2000000000

Post-processing: Listing first 45 summaries

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4: /cgn2_6/ptodata/1/ina/6B.COMB.seq.*
5: /cgn2_6/ptodata/1/ina/PTCUTS.COMB.seq.*
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Pred. No. is the number of results predicted by chance to have a score greater than or equal to the score of the result being printed, and is derived by analysis of the total score distribution.

SUMMARIES

Result No.	Score	Query Match	Length	ID	Description
1	59	100.0	59	US-08-864-357F-8	Sequence 8, Appl
2	37	62.7	59	US-08-864-357F-11	Sequence 11, Appl
3	17	28.8	14736	US-08-961-527F-171	Sequence 171, Appl
4	16	27.1	60	US-08-864-357F-10	Sequence 10, Appl
5	16	27.1	603	US-09-252-991A-3153	Sequence 3153, App
6	16	27.1	933	US-09-252-991A-8153	Sequence 815, App
7	16	27.1	2142	US-09-252-991A-869	Sequence 869, App
8	15	25.4	309	US-09-489-039A-1471	Sequence 1471, App
9	15	25.4	342	US-09-252-991A-3202	Sequence 3202, App
10	15	25.4	369	US-09-489-039A-1498	Sequence 1498, App
11	15	25.4	484	US-09-270-767-7800	Sequence 7800, App
12	15	25.4	484	US-09-270-767-23082	Sequence 23082, App
13	15	25.4	516	US-09-270-767-1781	Sequence 1781, App
14	15	25.4	516	US-09-270-767-17063	Sequence 17063, App
15	15	25.4	525	US-09-489-039A-1476	Sequence 1476, App
16	15	25.4	768	US-09-270-767-29574	Sequence 29574, App
17	15	25.4	807	US-09-248-796A-4201	Sequence 4201, App
18	15	25.4	987	US-09-252-991A-3011	Sequence 3011, App
19	15	25.4	1092	US-09-543-681A-3173	Sequence 3173, App
20	15	25.4	1296	US-09-270-767-13572	Sequence 13572, App
21	15	25.4	1534	US-08-629-643A-4	Sequence 4, Appl
22	15	25.4	1534	US-09-155-884-4	Sequence 4, Appl
23	15	25.4	1605	US-09-807-802A-16	Sequence 16, Appl
24	15	25.4	1632	US-09-252-991A-3113	Sequence 3113, App
25	15	25.4	1743	US-09-252-991A-2863	Sequence 2863, App
26	15	25.4	1800	US-09-807-802A-14	Sequence 14, Appl
27	15	25.4	2211	US-09-807-802A-12	Sequence 12, Appl

28	15	25.4	3000	US-09-705-267A-18	Sequence 18, Appl
29	15	25.4	4718	US-09-807-802A-1	Sequence 1, Appl
30	15	25.4	6727	US-08-629-643A-5	Sequence 5, Appl
31	15	25.4	6727	US-09-280-799-1	Sequence 1, Appl
32	15	25.4	6727	US-09-155-884-5	Sequence 5, Appl
33	15	25.4	7447	US-10-216-870-11	Sequence 11, Appl
34	15	25.4	9347	US-10-204-708-35	Sequence 35, Appl
35	15	25.4	10480	US-09-732-615-13	Sequence 13, Appl
36	15	25.4	10480	US-10-273-051-13	Sequence 13, Appl
37	15	25.4	35100	US-08-770-379-18	Sequence 18, Appl
38	15	25.4	35100	US-08-757-669A-18	Sequence 18, Appl
39	15	25.4	35100	US-09-230-571A-18	Sequence 18, Appl
40	15	25.4	786431	US-09-751-389-3	Sequence 3, Appl
41	15	25.4	1830121	US-09-557-884-1	Sequence 1, Appl
42	15	25.4	1830121	US-09-643-990A-1	Sequence 1, Appl
43	15	25.4	1830121	US-10-329-960-1	Sequence 1, Appl
44	15	23.7	47	US-09-671-317-817	Sequence 817, App
45	14	23.7	133	PCT-US96-05611A-7	Sequence 7, Appl

ALIGNMENTS

```

RESULT 1
US-08-864-357F-8
Sequence 8, Application US/08864357F
Patent No. 6255281
GENERAL INFORMATION:
APPLICANT: Claragen, Inc. & NIH
TITLE OF INVENTION: Use of Recombinant Human Uteroglobulin in Treatment of Inflammato
FILE REFERENCE: 116142/2
CURRENT APPLICATION NUMBER: US/08/864,357F
CURRENT FILING DATE: 1997-05-28
NUMBER OF SEQ ID NOS: 22
SOFTWARE: PatentIn version 3.0
SEQ ID NO: 8
LENGTH: 59
TYPE: DNA
ORGANISM: artificial
FEATURE:
OTHER INFORMATION: primer sequence
US-08-864-357F-8

Query Match      100.0%; Score 59; DB 3; Length 59;
Best Local Similarity 100.0%; Pred. No. 5.5e-24;
Matches 59; Conservative 0; Mismatches 0; Indels 0; Gaps 0;

Cy      1 CAGCTGAAGAAGTGGTTGACACCGCCGACAGAAACGGGTGAATCATCAATAACTG 59
Db      1 CAGCTGAAGAAGTGGTTGACACCGCCGACAGAAACGGGTGAATCATCAATAACTG 59

RESULT 2
US-08-864-357F-11/C
Sequence 11, Application US/08864357F
Patent No. 6255281
GENERAL INFORMATION:
APPLICANT: Claragen, Inc. & NIH
TITLE OF INVENTION: Use of Recombinant Human Uteroglobulin in Treatment of Inflammato
FILE REFERENCE: 116142/2
CURRENT APPLICATION NUMBER: US/08/864,357F
CURRENT FILING DATE: 1997-05-28
NUMBER OF SEQ ID NOS: 22
SOFTWARE: PatentIn version 3.0
SEQ ID NO: 11
LENGTH: 59
TYPE: DNA
ORGANISM: artificial
FEATURE:
OTHER INFORMATION: primer sequence
US-08-864-357F-11

```

Query Match 62.7%; Score 37; DB 3; Length 59;
Best Local Similarity 100.0%; Pred. No. 1.2e-11;
Matches 37; Conservative 0; Mismatches 0; Indels 0; Gaps 0;

CY 1 CAGCTGAAGAACTGGTTGACCCCTGCCGAGAAAC 37
DB 37 CAGCTGAAGAACTGGTTGACCCCTGCCGAGAAAC 1

RESULT 3

US-08-961-527-171/C
Sequence 171, Application US/08961527
Patent No. 6420135

GENERAL INFORMATION:
APPLICANT: Charles Kunsch
TITLE OF INVENTION: Streptococcus pneumoniae Polynucleotides and Sequences
NUMBER OF SEQUENCES: 391
CORRESPONDENCE ADDRESS:
ADDRESSEE: Human Genome Sciences, Inc.
STREET: 9410 Key West Avenue
CITY: Rockville
STATE: Maryland
COUNTRY: USA
ZIP: 20850
COMPUTER READABLE FORM:
MEDIUM TYPE: Diskette, 3.50 inch, 1.4MB storage
COMPUTER: HP Vectra 486/33
OPERATING SYSTEM: MSDOS version 6.2
SOFTWARE: ASCII text
CURRENT APPLICATION DATA:
APPLICATION NUMBER: US/08/961,527
FILING DATE:
CLASSIFICATION: 424
PRIOR APPLICATION DATA:
APPLICATION NUMBER:
FILING DATE:
ATTORNEY/AGENT INFORMATION:
NAME: Brookes, A. Anders
REGISTRATION NUMBER: 36,373
REFERENCE/DOCKET NUMBER: PB340P1
TELECOMMUNICATION INFORMATION:
TELEPHONE: (301) 309-8504
TELEFAX: (301) 309-8512
INFORMATION FOR SEQ ID NO: 171:
SEQUENCE CHARACTERISTICS:
LENGTH: 14736 base pairs
TYPE: nucleic acid
STRANDEDNESS: double
TOPOLOGY: linear
US-08-961-527-171

Query Match 28.8%; Score 17; DB 4; Length 14736;
Best Local Similarity 100.0%; Pred. No. 2.7;
Matches 17; Conservative 0; Mismatches 0; Indels 0; Gaps 0;

CY 42 TGAATCATCATTAAC 58
DB 13144 TGAATCATCATTAAC 13128

RESULT 4
US-08-864-357F-10/C
Sequence 10, Application US/08864357F
Patent No. 6255281

GENERAL INFORMATION:
APPLICANT: Clargen, Inc. & NIH
TITLE OF INVENTION: Use of Recombinant Human Uteroglobulin in Treatment of Inflammato
TITLE OF INVENTION: Fibrinolytic Conditions
FILE REFERENCE: 116142/2
CURRENT APPLICATION NUMBER: US/08/864,357F
CURRENT FILING DATE: 1997-05-28
NUMBER OF SEQ ID NOS: 22

SOFTWARE: PatentIn version 3.0

SEQ ID NO 10
LENGTH: 60
TYPE: DNA
ORGANISM: artificial
FEATURE:
OTHER INFORMATION: primer sequence
US-08-864-357F-10

Query Match 27.1%; Score 16; DB 3; Length 60;
Best Local Similarity 100.0%; Pred. No. 7.6;
Matches 16; Conservative 0; Mismatches 0; Indels 0; Gaps 0;

CY 38 CCGGTGAATCATCAT 53
DB 60 CCGGTGAATCATCAT 45

RESULT 5

US-09-252-991A-3153/C
Sequence 3153, Application US/09252991A
Patent No. 6551795

GENERAL INFORMATION:
APPLICANT: Marc J. Rubenfield et al.
TITLE OF INVENTION: NUCLEIC ACID AND AMINO ACID SEQUENCES RELATING TO PSEUDOMONAS
FILE REFERENCE: 107196.136
CURRENT APPLICATION NUMBER: US/09/252,991A
CURRENT FILING DATE: 1999-02-18
PRIOR APPLICATION NUMBER: US 60/074,788
PRIOR FILING DATE: 1998-02-18
PRIOR APPLICATION NUMBER: US 60/094,190
PRIOR FILING DATE: 1998-07-27
NUMBER OF SEQ ID NOS: 33142
SEQ ID NO 3153
LENGTH: 603
TYPE: DNA
ORGANISM: Pseudomonas aeruginosa
US-09-252-991A-3153

Query Match 27.1%; Score 16; DB 4; Length 603;
Best Local Similarity 100.0%; Pred. No. 8.4;
Matches 16; Conservative 0; Mismatches 0; Indels 0; Gaps 0;

CY 24 CCGTCCGCGAGAAACCG 39
DB 465 CCGTCCGCGAGAAACCG 450

RESULT 6

US-09-252-991A-810
Sequence 810, Application US/09252991A
Patent No. 6551795

GENERAL INFORMATION:
APPLICANT: Marc J. Rubenfield et al.
TITLE OF INVENTION: NUCLEIC ACID AND AMINO ACID SEQUENCES RELATING TO PSEUDOMONAS
FILE REFERENCE: 107196.136
CURRENT APPLICATION NUMBER: US/09/252,991A
CURRENT FILING DATE: 1999-02-18
PRIOR APPLICATION NUMBER: US 60/074,788
PRIOR FILING DATE: 1998-02-18
PRIOR APPLICATION NUMBER: US 60/094,190
PRIOR FILING DATE: 1998-07-27
NUMBER OF SEQ ID NOS: 33142
SEQ ID NO 810
LENGTH: 933
TYPE: DNA
ORGANISM: Pseudomonas aeruginosa
US-09-252-991A-810

Query Match 27.1%; Score 16; DB 4; Length 933;
Best Local Similarity 100.0%; Pred. No. 8.5;

Matches 16; Conservative 0; Mismatches 0; Indels 0; Gaps 0;
QY 18 TGACACCTGGCGGAG 33
Db 564 TGACACCTGGCGGAG 579

RESULT 7
US-09-252-991A-869/c
Sequence 869, Application US/09252991A
Patent No. 6551795
GENERAL INFORMATION:
APPLICANT: Marc J. Rubenfield et al.
TITLE OF INVENTION: NUCLEIC ACID AND AMINO ACID SEQUENCES RELATING TO PSEUDOMONAS
FILE REFERENCE: 107196.136
CURRENT APPLICATION NUMBER: US/09/252,991A
CURRENT FILING DATE: 1999-02-18
PRIOR APPLICATION NUMBER: US 60/074,788
PRIOR FILING DATE: 1998-02-18
PRIOR APPLICATION NUMBER: US 60/094,190
PRIOR FILING DATE: 1998-07-27
NUMBER OF SEQ ID NOS: 33142
SEQ ID NO 869
LENGTH: 2142
TYPE: DNA
ORGANISM: Pseudomonas aeruginosa
US-09-252-991A-869

Query Match 27.1%; Score 16; DB 4; Length 2142;
Best Local Similarity 100.0%; Pred. No. 8.9;
Matches 16; Conservative 0; Mismatches 0; Indels 0; Gaps 0;
QY 18 TGACACCTGGCGGAG 33
Db 1639 TGACACCTGGCGGAG 1624

RESULT 8
US-09-489-039A-1471
Sequence 1471, Application US/09489039A
Patent No. 6610836
GENERAL INFORMATION:
APPLICANT: Gary Breton et al.
TITLE OF INVENTION: NUCLEIC ACID AND AMINO ACID SEQUENCES RELATING TO KLEBSIELLA
FILE REFERENCE: 2709.2004001
CURRENT APPLICATION NUMBER: US/09/489,039A
CURRENT FILING DATE: 2000-01-27
PRIOR APPLICATION NUMBER: US 60/117,747
PRIOR FILING DATE: 1999-01-29
NUMBER OF SEQ ID NOS: 14342
SEQ ID NO 1471
LENGTH: 309
TYPE: DNA
ORGANISM: Klebsiella pneumoniae
US-09-489-039A-1471

Query Match 25.4%; Score 15; DB 4; Length 309;
Best Local Similarity 100.0%; Pred. No. 30;
Matches 15; Conservative 0; Mismatches 0; Indels 0; Gaps 0;
QY 3 GCTGAAGAACTGGT 17
Db 237 GCTGAAGAACTGGT 251

RESULT 9
US-09-252-991A-3202
Sequence 3202, Application US/09252991A
Patent No. 6551795
GENERAL INFORMATION:
APPLICANT: Marc J. Rubenfield et al.

TITLE OF INVENTION: NUCLEIC ACID AND AMINO ACID SEQUENCES RELATING TO PSEUDOMONAS
FILE REFERENCE: 107196.136
CURRENT APPLICATION NUMBER: US/09/252,991A
CURRENT FILING DATE: 1999-02-18
PRIOR APPLICATION NUMBER: US 60/074,788
PRIOR FILING DATE: 1998-02-18
PRIOR APPLICATION NUMBER: US 60/094,190
PRIOR FILING DATE: 1998-07-27
NUMBER OF SEQ ID NOS: 33142
SEQ ID NO 3202
LENGTH: 342
TYPE: DNA
ORGANISM: Pseudomonas aeruginosa
US-09-252-991A-3202

Query Match 25.4%; Score 15; DB 4; Length 342;
Best Local Similarity 100.0%; Pred. No. 30;
Matches 15; Conservative 0; Mismatches 0; Indels 0; Gaps 0;
QY 21 CACCTGGCGGAGAA 35
Db 2 CACCTGGCGGAGAA 16

RESULT 10
US-09-489-039A-1498/c
Sequence 1498, Application US/09489039A
Patent No. 6610836
GENERAL INFORMATION:
APPLICANT: Gary Breton et al.
TITLE OF INVENTION: NUCLEIC ACID AND AMINO ACID SEQUENCES RELATING TO KLEBSIELLA
FILE REFERENCE: 2709.2004001
CURRENT APPLICATION NUMBER: US/09/489,039A
CURRENT FILING DATE: 2000-01-27
PRIOR APPLICATION NUMBER: US 60/117,747
PRIOR FILING DATE: 1999-01-29
NUMBER OF SEQ ID NOS: 14342
SEQ ID NO 1498
LENGTH: 369
TYPE: DNA
ORGANISM: Klebsiella pneumoniae
US-09-489-039A-1498

Query Match 25.4%; Score 15; DB 4; Length 369;
Best Local Similarity 100.0%; Pred. No. 30;
Matches 15; Conservative 0; Mismatches 0; Indels 0; Gaps 0;
QY 3 GCTGAAGAACTGGT 17
Db 124 GCTGAAGAACTGGT 110

RESULT 11
US-09-270-767-7800/c
Sequence 7800, Application US/09270767
Patent No. 6703491
GENERAL INFORMATION:
APPLICANT: Homburger et al.
TITLE OF INVENTION: Nucleic acids and proteins of Drosophila melanogaster
FILE REFERENCE: File Reference: 7326-094
CURRENT APPLICATION NUMBER: US/09/270,767
CURRENT FILING DATE: 1999-03-17
NUMBER OF SEQ ID NOS: 62517
SOFTWARE: PatentIn Ver. 2.0
SEQ ID NO 7800
LENGTH: 484
TYPE: DNA
ORGANISM: Drosophila melanogaster
US-09-270-767-7800

Query Match 25.4%; Score 15; DB 4; Length 484;

Best Local Similarity 100.0%; Pred. No. 30;
Matches 15; Conservative 0; Mismatches 0; Indels 0; Gaps 0;
QY 1 CAGCTGAAGAACTG 15
Db 366 CAGCTGAAGAACTG 352

RESULT 12
US-09-270-767-23082/c
Sequence 23082, Application US/09270767
Patent No. 6703491
GENERAL INFORMATION:
APPLICANT: Homburger et al.
TITLE OF INVENTION: Nucleic acids and proteins of *Drosophila melanogaster*
FILE REFERENCE: File Reference: 7326-094
CURRENT FILING DATE: 1999-03-17
NUMBER OF SEQ ID NOS: 62517
SOFTWARE: Patentia Ver. 2.0
SEQ ID NO 23082
LENGTH: 484
TYPE: DNA
ORGANISM: *Drosophila melanogaster*
US-09-270-767-23082

Query Match 25.4%; Score 15; DB 4; Length 484;
Best Local Similarity 100.0%; Pred. No. 30;
Matches 15; Conservative 0; Mismatches 0; Indels 0; Gaps 0;

QY 1 CAGCTGAAGAACTG 15
Db 366 CAGCTGAAGAACTG 352

RESULT 13
US-09-270-767-1781/c
Sequence 1781, Application US/09270767
Patent No. 6703491
GENERAL INFORMATION:
APPLICANT: Homburger et al.
TITLE OF INVENTION: Nucleic acids and proteins of *Drosophila melanogaster*
FILE REFERENCE: File Reference: 7326-094
CURRENT FILING DATE: 1999-03-17
NUMBER OF SEQ ID NOS: 62517
SOFTWARE: Patentia Ver. 2.0
SEQ ID NO 1781
LENGTH: 516
TYPE: DNA
ORGANISM: *Drosophila melanogaster*
US-09-270-767-1781

Query Match 25.4%; Score 15; DB 4; Length 516;
Best Local Similarity 100.0%; Pred. No. 30;
Matches 15; Conservative 0; Mismatches 0; Indels 0; Gaps 0;

QY 1 CAGCTGAAGAACTG 15
Db 432 CAGCTGAAGAACTG 418

RESULT 14
US-09-270-767-17063/c
Sequence 17063, Application US/09270767
Patent No. 6703491
GENERAL INFORMATION:
APPLICANT: Homburger et al.
TITLE OF INVENTION: Nucleic acids and proteins of *Drosophila melanogaster*
FILE REFERENCE: File Reference: 7326-094
CURRENT FILING DATE: 1999-03-17
NUMBER OF SEQ ID NOS: 62517

SOFTWARE: Patentia Ver. 2.0
SEQ ID NO 17063
LENGTH: 516
TYPE: DNA
ORGANISM: *Drosophila melanogaster*
US-09-270-767-17063

Query Match 25.4%; Score 15; DB 4; Length 516;
Best Local Similarity 100.0%; Pred. No. 30;
Matches 15; Conservative 0; Mismatches 0; Indels 0; Gaps 0;

QY 1 CAGCTGAAGAACTG 15
Db 432 CAGCTGAAGAACTG 418

RESULT 15
US-09-489-039A-1476/c
Sequence 1476, Application US/09489039A
Patent No. 6610836
GENERAL INFORMATION:
APPLICANT: Gary Bretton et al.
TITLE OF INVENTION: NUCLEIC ACID AND AMINO ACID SEQUENCES RELATING TO *KLEBSIELLA*
FILE REFERENCE: 2703.2004001
CURRENT FILING DATE: 2000-01-27
PRIOR APPLICATION NUMBER: US 60/117,747
PRIOR FILING DATE: 1999-01-29
NUMBER OF SEQ ID NOS: 14342
SEQ ID NO 1476
LENGTH: 525
TYPE: DNA
ORGANISM: *Klebsiella pneumoniae*
US-09-489-039A-1476

Query Match 25.4%; Score 15; DB 4; Length 525;
Best Local Similarity 100.0%; Pred. No. 30;
Matches 15; Conservative 0; Mismatches 0; Indels 0; Gaps 0;

QY 3 GCTGAAGAACTGCT 17
Db 444 GCTGAAGAACTGCT 430

Search completed: December 23, 2004, 01:33:42
Job time : 53.5204 secs

GenCore version 5.1.6
Copyright (c) 1993 - 2004 CompuGen Ltd.

OM nucleic - nucleic search, using sw model

Run on: December 22, 2004, 23:36:53 ; Search time 827.066 Seconds
(without alignments)
397.214 Million cell updates/sec

Title: US-09-898-616A-3

Perfect score: 59
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Scoring table: OLIGO NUC
Gapop 60.0 , Gapext 60.0

Searched: 4105333 seqs, 2784095677 residues

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Minimum DB seq length: 0
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Pred. No. is the number of results predicted by chance to have a score greater than or equal to the score of the result being printed, and is derived by analysis of the total score distribution.

SUMMARIES

Result No.	Score	Query Match	Length	ID	Description
1	59	100.0	59	9	US-09-861-688-8
2	59	100.0	59	10	US-09-898-616A-3
3	59	100.0	59	15	US-10-187-498A-3
4	59	100.0	59	16	US-10-647-371-7
5	37	62.7	59	9	US-09-861-688-11
6	37	62.7	59	10	US-09-898-616A-6
7	37	62.7	59	15	US-10-187-498A-6
8	37	62.7	59	16	US-10-647-371-10
9	19	32.2	757	17	US-10-437-963-89235
10	18	30.5	1152	17	US-10-437-963-89235
11	18	30.5	1152	17	US-10-437-963-89235
12	18	30.5	2004	9	US-09-867-576-235

13	18	30.5	2134	17	US-10-437-963-74313	Sequence 74313, A
14	17	28.8	195	9	US-09-974-300-3436	Sequence 3436, Ap
15	17	28.8	520	13	US-10-027-632-51350	Sequence 51350, A
16	17	28.8	520	15	US-10-027-632-51350	Sequence 51350, A
17	17	28.8	549	13	US-10-027-632-71225	Sequence 71225, A
18	17	28.8	549	13	US-10-027-632-71225	Sequence 71225, A
19	17	28.8	549	13	US-10-027-632-71225	Sequence 71225, A
20	17	28.8	549	15	US-10-027-632-71225	Sequence 71225, A
21	17	28.8	868	13	US-10-027-632-72285	Sequence 72285, A
22	17	28.8	868	15	US-10-027-632-72285	Sequence 72285, A
23	17	28.8	993	15	US-10-146-742-317	Sequence 317, App
24	17	28.8	993	16	US-10-440-503-317	Sequence 317, App
25	17	28.8	993	16	US-10-440-503-317	Sequence 317, App
26	17	28.8	993	16	US-10-440-503-317	Sequence 317, App
27	17	28.8	993	16	US-10-440-503-317	Sequence 317, App
28	17	28.8	1000	16	US-10-425-114-16587	Sequence 16587, A
29	17	28.8	1130	18	US-10-425-115-15378	Sequence 15378, A
30	17	28.8	14736	8	US-08-961-527-171	Sequence 171, App
31	17	28.8	14736	16	US-10-158-844-171	Sequence 171, App
32	17	28.8	59475	17	US-10-332-696-166	Sequence 166, App
33	16	27.1	60	10	US-09-861-688-10	Sequence 10, Appl
34	16	27.1	60	10	US-09-898-616A-5	Sequence 5, Appl
35	16	27.1	60	15	US-10-187-498A-5	Sequence 5, Appl
36	16	27.1	60	16	US-10-647-371-9	Sequence 9, Appl
37	16	27.1	311	10	US-09-764-891-1420	Sequence 1420, Ap
38	16	27.1	373	16	US-10-424-599-11255	Sequence 11255, A
39	16	27.1	377	9	US-09-983-965-5306	Sequence 5306, Ap
40	16	27.1	455	18	US-10-425-115-115741	Sequence 115741, A
41	16	27.1	547	16	US-10-424-599-35982	Sequence 35982, A
42	16	27.1	639	18	US-10-320-797-2026	Sequence 2026, Ap
43	16	27.1	659	16	US-10-653-047-581	Sequence 581, App
44	16	27.1	667	13	US-10-005-057A-34	Sequence 34, Appl
45	16	27.1	667	16	US-10-675-072A-37	Sequence 37, Appl

ALIGNMENTS

RESULT 1

US-09-861-688-8

Sequence 8, Appli

Patent No. US20020173460A1

GENERAL INFORMATION:

APPLICANT: Clargen, Inc. & NIH

TITLE OF INVENTION: Use of Recombinant Human Uteroglobin in Treatment of

TITLE OF INVENTION: Inflammatory and

TITLE OF INVENTION: Fibrotic Conditions

FILE REFERENCE: 116142/2

CURRENT APPLICATION NUMBER: US/09/861,688

CURRENT FILING DATE: 2001-05-21

PRIOR APPLICATION NUMBER: 08/864,357

PRIOR FILING DATE: 1997-05-28

NUMBER OF SEQ ID NOS: 22

SOFTWARE: PatentIn version 3.0

SEQ ID NO 8

LENGTH: 59

TYPE: DNA

ORGANISM: Artificial Sequence

FEATURE:

OTHER INFORMATION: primer sequence

US-09-861-688-8

Query Match

Best Local Similarity 100.0%; Score 59; DB 9; Length 59;

Matches 59; Conservative 0; Mismatches 0; Indels 0; Gaps 0;

Qy

1 CAGCTGAAGAAGTGTGACACCCCTCCGAGAAACCGGTGATCATCAATAACTG 59

Db

1 CAGCTGAAGAAGTGTGACACCCCTCCGAGAAACCGGTGATCAATCAATAACTG 59

RESULT 2

US-09-898-616A-3

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Sequence 3, Application US/09898616A
Publication No. US20030109429A1
GENERAL INFORMATION:
APPLICANT: Clargen Inc.
APPLICANT: Pilon, Aprile L
APPLICANT: Welch, Richard W
TITLE OF INVENTION: Method for the Production of Purified rhug for the Treatment of
FILE REFERENCE: 116142/00170
CURRENT APPLICATION NUMBER: US/09/898,616A
PRIOR FILING DATE: 2002-10-15
PRIOR APPLICATION NUMBER: US 08/864,357
NUMBER OF SEQ ID NOS: 10
SOFTWARE: PatentIn version 3.1
SEQ ID NO 3
LENGTH: 59
TYPE: DNA
ORGANISM: Artificial Sequence
FEATURE:
OTHER INFORMATION: Original amino acid sequence obtained from cDNA. Current submitte
FEATURE:
NAME/KEY: misc feature
OTHER INFORMATION: Original amino acid sequence obtained from cDNA. Current submitte
US-09-898-616A-3

Query Match 100.0%; Score 59; DB 10; Length 59;
Best Local Similarity 100.0%; Pred. No. 1.7e-23;
Matches 59; Conservative 0; Mismatches 0; Indels 0; Gaps 0;

Cy 1 CAGCTGAAGAACTGGTTGACACCCCTGCCGAGAAACCGCGTGAATCATCAATAACTG 59
Db 1 CAGCTGAAGAACTGGTTGACACCCCTGCCGAGAAACCGCGTGAATCATCAATAACTG 59

RESULT 3
US-10-187-498A-3
Sequence 3, Application US/10187498A
Publication No. US20030207925A1
GENERAL INFORMATION:
APPLICANT: Clargen Inc.
APPLICANT: Pilon, Aprile L
APPLICANT: Welch, Richard W
TITLE OF INVENTION: Method for the Production of Purified rhug for the Treatment of
FILE REFERENCE: 116142/00260
CURRENT APPLICATION NUMBER: US/10/187,498A
PRIOR FILING DATE: 2001-07-02
PRIOR APPLICATION NUMBER: US 08/864,357
NUMBER OF SEQ ID NOS: 10
SOFTWARE: PatentIn version 3.1
SEQ ID NO 3
LENGTH: 59
TYPE: DNA
ORGANISM: Artificial Sequence
FEATURE:
OTHER INFORMATION: Original amino acid sequence obtained from cDNA. Current submitte
FEATURE:
NAME/KEY: misc feature
OTHER INFORMATION: Original amino acid sequence obtained from cDNA. Current submitte
US-10-187-498A-3

Query Match 100.0%; Score 59; DB 15; Length 59;
Best Local Similarity 100.0%; Pred. No. 1.7e-23;
Matches 59; Conservative 0; Mismatches 0; Indels 0; Gaps 0;

Cy 1 CAGCTGAAGAACTGGTTGACACCCCTGCCGAGAAACCGCGTGAATCATCAATAACTG 59
Db 1 CAGCTGAAGAACTGGTTGACACCCCTGCCGAGAAACCGCGTGAATCATCAATAACTG 59
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Db 1 CAGCTGAAGAACTGGTTGACACCCCTGCCGAGAAACCGCGTGAATCATCAATAACTG 59

RESULT 4
US-10-647-371-7
Sequence 7, Application US/10647371
Publication No. US20040047857A1
GENERAL INFORMATION:
APPLICANT: Clargen, Inc. & NIH
TITLE OF INVENTION: Use of Recombinant Human Uteroglobulin in Treatment of Inflammatory
FILE REFERENCE: 116142-85
CURRENT APPLICATION NUMBER: US/10/647,371
PRIOR FILING DATE: 2003-08-25
PRIOR APPLICATION NUMBER: 09/549,926
NUMBER OF SEQ ID NOS: 12
SOFTWARE: PatentIn version 3.2
SEQ ID NO 7
LENGTH: 59
TYPE: DNA
ORGANISM: Artificial
FEATURE:
OTHER INFORMATION: Description of Artificial Sequence: primer sequence
US-10-647-371-7

Query Match 100.0%; Score 59; DB 16; Length 59;
Best Local Similarity 100.0%; Pred. No. 1.7e-23;
Matches 59; Conservative 0; Mismatches 0; Indels 0; Gaps 0;

Cy 1 CAGCTGAAGAACTGGTTGACACCCCTGCCGAGAAACCGCGTGAATCATCAATAACTG 59
Db 1 CAGCTGAAGAACTGGTTGACACCCCTGCCGAGAAACCGCGTGAATCATCAATAACTG 59

RESULT 5
US-09-861-688-11/c
Sequence 11, Application US/09861688
Patent No. US20020173460A1
GENERAL INFORMATION:
APPLICANT: Clargen, Inc. & NIH
TITLE OF INVENTION: Use of Recombinant Human Uteroglobulin in Treatment of
FILE REFERENCE: 116142/2
CURRENT APPLICATION NUMBER: US/09/861,688
PRIOR FILING DATE: 2001-05-21
PRIOR APPLICATION NUMBER: 08/864,357
NUMBER OF SEQ ID NOS: 22
SOFTWARE: PatentIn version 3.0
SEQ ID NO 11
LENGTH: 59
TYPE: DNA
ORGANISM: Artificial Sequence
FEATURE:
OTHER INFORMATION: primer sequence
US-09-861-688-11

Query Match 62.7%; Score 37; DB 9; Length 59;
Best Local Similarity 100.0%; Pred. No. 5.4e-11;
Matches 37; Conservative 0; Mismatches 0; Indels 0; Gaps 0;

Cy 1 CAGCTGAAGAACTGGTTGACACCCCTGCCGAGAAAC 37
Db 37 CAGCTGAAGAACTGGTTGACACCCCTGCCGAGAAAC 1

RESULT 6
US-09-898-616A-6/c
Sequence 6, Application US/09898616A
Publication No. US20030109429A1
GENERAL INFORMATION:
```



```
APPLICANT: Clargen Inc.
APPLICANT: Pilon, Aprile L
APPLICANT: Welch, Richard W
TITLE OF INVENTION: Method for the Production of Purified rhug for the Treatment of
FILE REFERENCE: 116142/00170
CURRENT APPLICATION NUMBER: US/09/898,616A
CURRENT FILING DATE: 2002-10-15
PRIOR APPLICATION NUMBER: US 08/864,357
PRIOR FILING DATE: 1997-05-28
NUMBER OF SEQ ID NOS: 10
SOFTWARE: Patent version 3.1
SEQ ID NO 6
LENGTH: 59
TYPE: DNA
ORGANISM: Artificial Sequence
FEATURE:
OTHER INFORMATION: Original amino acid sequence obtained from cDNA. Current submitte
OTHER INFORMATION: d sequence maximized for expression in E. coli.
NAME/KEY: misc.feature
OTHER INFORMATION: Original amino acid sequence obtained from cDNA. Current submitte
OTHER INFORMATION: d sequence maximized for expression in E. coli.
US-09-898-616A-6
```

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Query Match 62.7%; Score 37; DB 10; Length 59;
Best Local Similarity 100.0%; Pred. No. 5,4e-11;
Matches 37; Conservative 0; Mismatches 0; Indels 0; Gaps 0;
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QY 1 CAGCTGAAGAAACTGTTGACACCCCTGCGCAGAAAC 37
DB 37 CAGCTGAAGAAACTGTTGACACCCCTGCGCAGAAAC 1
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```
RESULT 7
US-10-187-498A-6/c
Sequence 6, Application US/10187498A
Publication No. US2003020795A1
GENERAL INFORMATION:
APPLICANT: Clargen Inc.
APPLICANT: Pilon, Aprile L
APPLICANT: Welch, Richard W
TITLE OF INVENTION: Method for the Production of Purified rhug for the Treatment of
FILE REFERENCE: 116142/00260
CURRENT APPLICATION NUMBER: US/10/187,498A
CURRENT FILING DATE: 2001-07-02
PRIOR APPLICATION NUMBER: US 08/864,357
PRIOR FILING DATE: 1997-05-28
NUMBER OF SEQ ID NOS: 10
SOFTWARE: Patent version 3.1
SEQ ID NO 6
LENGTH: 59
TYPE: DNA
ORGANISM: Artificial Sequence
FEATURE:
OTHER INFORMATION: Original amino acid sequence obtained from cDNA. Current submitte
OTHER INFORMATION: d sequence maximized for expression in E. coli.
NAME/KEY: misc.feature
OTHER INFORMATION: Original amino acid sequence obtained from cDNA. Current submitte
OTHER INFORMATION: d sequence maximized for expression in E. coli.
US-10-187-498A-6
```

```
Query Match 62.7%; Score 37; DB 15; Length 59;
Best Local Similarity 100.0%; Pred. No. 5,4e-11;
Matches 37; Conservative 0; Mismatches 0; Indels 0; Gaps 0;
```

```
QY 1 CAGCTGAAGAAACTGTTGACACCCCTGCGCAGAAAC 37
DB 37 CAGCTGAAGAAACTGTTGACACCCCTGCGCAGAAAC 1
```

```
RESULT 8
US-10-647-371-10/c
Sequence 10, Application US/10647371
Publication No. US20040047857A1
GENERAL INFORMATION:
APPLICANT: Clargen, Inc. & NIH
TITLE OF INVENTION: Use of Recombinant Human Uteroglobulin in Treatment of Inflammatory
FILE REFERENCE: 116142-85
CURRENT APPLICATION NUMBER: US/10/647,371
CURRENT FILING DATE: 2003-08-25
PRIOR APPLICATION NUMBER: 09/549,926
PRIOR FILING DATE: 2000-04-14
NUMBER OF SEQ ID NOS: 12
SOFTWARE: Patent version 3.2
SEQ ID NO 10
LENGTH: 59
TYPE: DNA
ORGANISM: Artificial
FEATURE:
OTHER INFORMATION: Description of Artificial Sequence: primer sequence
US-10-647-371-10
```

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Query Match 62.7%; Score 37; DB 16; Length 59;
Best Local Similarity 100.0%; Pred. No. 5,4e-11;
Matches 37; Conservative 0; Mismatches 0; Indels 0; Gaps 0;
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QY 1 CAGCTGAAGAAACTGTTGACACCCCTGCGCAGAAAC 37
DB 37 CAGCTGAAGAAACTGTTGACACCCCTGCGCAGAAAC 1
```

```
RESULT 9
US-10-437-963-89235
Sequence 89235, Application US/10437963
Publication No. US20040123343A1
GENERAL INFORMATION:
APPLICANT: La Rosa, Thomas J.
APPLICANT: Kovalic, David K.
APPLICANT: Zhou, Yihua
APPLICANT: Cao, Yongwei
APPLICANT: Wu, Wei
APPLICANT: Boukharov, Andrey A.
APPLICANT: Barbazuk, Brad
APPLICANT: Li, Ping
TITLE OF INVENTION: Rice Nucleic Acid Molecules and Other Molecules Associated With
FILE REFERENCE: 38-21(53221)B
CURRENT APPLICATION NUMBER: US/10/437,963
CURRENT FILING DATE: 2003-05-14
NUMBER OF SEQ ID NOS: 204966
SEQ ID NO 89235
LENGTH: 757
TYPE: DNA
ORGANISM: Oryza sativa
FEATURE:
OTHER INFORMATION: Clone ID: PAT_MRT4530_88009C.1
US-10-437-963-89235
```

```
Query Match 32.2%; Score 19; DB 17; Length 757;
Best Local Similarity 100.0%; Pred. No. 1;
Matches 19; Conservative 0; Mismatches 0; Indels 0; Gaps 0;
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```
QY 2 AGCTGAAGAAACTGTTGA 20
DB 207 AGCTGAAGAAACTGTTGA 225
```

```
RESULT 10
US-10-437-963-65306/c
Sequence 65306, Application US/10437963
Publication No. US20040123343A1
GENERAL INFORMATION:
```

APPLICANT: La Rosa, Thomas J.
APPLICANT: Kovalic, David K.
APPLICANT: Zhou, Yihua
APPLICANT: Cao, Yongwei
APPLICANT: Wu, Wei
APPLICANT: Boukharov, Andrey A.
APPLICANT: Barbazuk, Brad
APPLICANT: Li, Ping
TITLE OF INVENTION: Rice Nucleic Acid Molecules and Other Molecules Associated With
FILE REFERENCE: 38-21(53221)B
CURRENT APPLICATION NUMBER: US/10/437,963
CURRENT FILING DATE: 2003-05-14
NUMBER OF SEQ ID NOS: 204966
SEQ ID NO 65306
LENGTH: 490
TYPE: DNA
ORGANISM: Oryza sativa
FEATURE:
OTHER INFORMATION: Clone ID: PAT_MRT4530_66367C.1
US-10-437-963-65306

Query Match 30.5%; Score 18; DB 17; Length 490;
Best Local Similarity 100.0%; Pred. No. 3.8;
Matches 18; Conservative 0; Mismatches 0; Indels 0; Gaps 0;

QY 13 CTGGTTGACACCCCTGCCG 30
DB 335 CTGGTTGACACCCCTGCCG 318

RESULT 11
US-10-437-963-65308/c
Sequence 65308, Application US/10437963
Publication No. US20040123343A1
GENERAL INFORMATION:
APPLICANT: La Rosa, Thomas J.
APPLICANT: Kovalic, David K.
APPLICANT: Zhou, Yihua
APPLICANT: Cao, Yongwei
APPLICANT: Wu, Wei
APPLICANT: Boukharov, Andrey A.
APPLICANT: Barbazuk, Brad
APPLICANT: Li, Ping
TITLE OF INVENTION: Rice Nucleic Acid Molecules and Other Molecules Associated With
FILE REFERENCE: 38-21(53221)B
CURRENT APPLICATION NUMBER: US/10/437,963
CURRENT FILING DATE: 2003-05-14
NUMBER OF SEQ ID NOS: 204966
SEQ ID NO 65308
LENGTH: 1152
TYPE: DNA
ORGANISM: Oryza sativa
FEATURE:
OTHER INFORMATION: Clone ID: PAT_MRT4530_66369C.1
US-10-437-963-65308

Query Match 30.5%; Score 18; DB 17; Length 1152;
Best Local Similarity 100.0%; Pred. No. 4;
Matches 18; Conservative 0; Mismatches 0; Indels 0; Gaps 0;

QY 13 CTGGTTGACACCCCTGCCG 30
DB 668 CTGGTTGACACCCCTGCCG 651

RESULT 12
US-09-887-576-235/c
Sequence 235, Application US/09887576
Patent No. US20020144047A1
GENERAL INFORMATION:
APPLICANT: Budworth, P.

APPLICANT: Brown, D.
APPLICANT: Chang, H.
APPLICANT: Zhu, T.
APPLICANT: Han, B.
APPLICANT: Wang, X.
APPLICANT: Cooper, Bret
TITLE OF INVENTION: Promoters for regulation of plant expression
FILE REFERENCE: 1360.001US1
CURRENT APPLICATION NUMBER: US/09/887,576
CURRENT FILING DATE: 2001-06-25
PRIOR APPLICATION NUMBER: US 60/213,848
PRIOR FILING DATE: 2000-06-23
PRIOR APPLICATION NUMBER: US 60/214,087
PRIOR FILING DATE: 2000-06-23
PRIOR APPLICATION NUMBER: US 60/258,692
PRIOR FILING DATE: 2000-12-29
NUMBER OF SEQ ID NOS: 875
SOFTWARE: FastSeq for Windows Version 4.0
SEQ ID NO 235
LENGTH: 2004
TYPE: DNA
ORGANISM: Arabidopsis thaliana
US-09-887-576-235

Query Match 30.5%; Score 18; DB 9; Length 2004;
Best Local Similarity 100.0%; Pred. No. 4.1;
Matches 18; Conservative 0; Mismatches 0; Indels 0; Gaps 0;

QY 30 GCAGAAACCGCGTGAATC 47
DB 389 GCAGAAACCGCGTGAATC 372

RESULT 13
US-10-437-963-74313
Sequence 74313, Application US/10437963
Publication No. US20040123343A1
GENERAL INFORMATION:
APPLICANT: La Rosa, Thomas J.
APPLICANT: Kovalic, David K.
APPLICANT: Zhou, Yihua
APPLICANT: Cao, Yongwei
APPLICANT: Wu, Wei
APPLICANT: Boukharov, Andrey A.
APPLICANT: Barbazuk, Brad
APPLICANT: Li, Ping
TITLE OF INVENTION: Rice Nucleic Acid Molecules and Other Molecules Associated With
FILE REFERENCE: 38-21(53221)B
CURRENT APPLICATION NUMBER: US/10/437,963
CURRENT FILING DATE: 2003-05-14
NUMBER OF SEQ ID NOS: 204966
SEQ ID NO 74313
LENGTH: 2134
TYPE: DNA
ORGANISM: Oryza sativa
FEATURE:
OTHER INFORMATION: Clone ID: PAT_MRT4530_74510C.1
US-10-437-963-74313

Query Match 30.5%; Score 18; DB 17; Length 2134;
Best Local Similarity 100.0%; Pred. No. 4.1;
Matches 18; Conservative 0; Mismatches 0; Indels 0; Gaps 0;

QY 3 GCTGAAGAACTGTTGA 20
DB 1728 GCTGAAGAACTGTTGA 1745

RESULT 14
US-09-874-300-3436/c
Sequence 3436, Application US/09974300
Patent No. US20020146721A1

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; GENERAL INFORMATION:
; APPLICANT: Berka, Randy M.
; APPLICANT: Clausen, Ib Groth
; TITLE OF INVENTION: Methods for Monitoring Multiple Gene
; FILE REFERENCE: 10085,500-US
; CURRENT APPLICATION NUMBER: US/09/974,300
; CURRENT FILING DATE: 2001-10-05
; PRIOR APPLICATION NUMBER: 09/680,598
; PRIOR FILING DATE: 2000-10-06
; PRIOR APPLICATION NUMBER: 60/279,526
; PRIOR FILING DATE: 2001-03-27
; NUMBER OF SEQ ID NOS: 8481
; SOFTWARE: FastSeq for Windows Version 4.0
; SEQ ID NO 3436
; LENGTH: 195
; TYPE: DNA
; ORGANISM: Bacillus licheniformis
US-09-974-300-3436

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Query Match      28.8%; Score 17; DB 9; Length 195;
Best Local Similarity 100.0%; Pred. No. 13;
Matches 17; Conservative 0; Mismatches 0; Indels 0; Gaps 0;

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QY      21 CACCTGCGCGAAGAC 37
Db      102 CACCTGCGCGAAGAC 86

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RESULT 15
US-10-027-632-51350/c
; Sequence 51350, Application US/10027632.
; Publication No. US20020198371A1
; GENERAL INFORMATION:
; APPLICANT: Wang, David G.
; TITLE OF INVENTION: Identification and Mapping of Single Nucleotide
; FILE REFERENCE: 108827.129
; CURRENT APPLICATION NUMBER: US/10/027,632
; CURRENT FILING DATE: 2002-04-30
; PRIOR APPLICATION NUMBER: US 60/218,006
; PRIOR FILING DATE: 2000-07-12
; PRIOR APPLICATION NUMBER: US 60/198,676
; PRIOR FILING DATE: 2000-04-20
; PRIOR APPLICATION NUMBER: US 60/193,483
; PRIOR FILING DATE: 2000-03-29
; PRIOR APPLICATION NUMBER: US 60/185,218
; PRIOR FILING DATE: 2000-02-24
; PRIOR APPLICATION NUMBER: US 60/167,363
; PRIOR FILING DATE: 1999-11-23
; PRIOR APPLICATION NUMBER: US 60/156,358
; PRIOR FILING DATE: 1999-09-28
; PRIOR APPLICATION NUMBER: US 60/146,002
; PRIOR FILING DATE: 1999-08-09
; NUMBER OF SEQ ID NOS: 325720
; SOFTWARE: FastSeq for Windows Version 4.0
; SEQ ID NO 51350
; LENGTH: 520
; TYPE: DNA
; ORGANISM: Human
US-10-027-632-51350

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Query Match      28.8%; Score 17; DB 13; Length 520;
Best Local Similarity 100.0%; Pred. No. 14;
Matches 17; Conservative 0; Mismatches 0; Indels 0; Gaps 0;

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QY      5 TGAAGAACTGGTTGAC 21
Db      106 TGAAGAACTGGTTGAC 90

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Search completed: December 23, 2004, 05:13:28
 Job time : 829.068 secs

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GenCore version 5.1.6
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OM nucleic - nucleic search, using sw model

Run on: December 22, 2004, 19:17:58 ; Search time 523.357 Seconds
(without alignments)
3343.258 Million cell updates/sec

Title: US-09-898-616a-4

Sequence: 1 atggaagaatgcctcagctcagctgtgcactaaag 37

Scoring table: OLIGO_NUC
Gapop 60.0 , Gapext 60.0

Searched: 4526729 seqs, 23644849745 residues

Word size : 0

Total number of hits satisfying chosen parameters: 9053458

Minimum DB seq length: 0
Maximum DB seq length: 2000000000

Post-processing: Listing first 45 summaries

Database : GenEmbl.*

1: gb_da:*
2: gb_hhg:*
3: gb_in:*
4: gb_om:*
5: gb_ov:*
6: gb_pat:*
7: gb_ph:*
8: gb_pl:*
9: gb_pr:*
10: gb_ro:*
11: gb_sts:*
12: gb_sy:*
13: gb_un:*
14: gb_vl:*

Pred. No. is the number of results predicted by chance to have a score greater than or equal to the score of the result being printed, and is derived by analysis of the total score distribution.

SUMMARIES

Result No.	Score	Query Match Length	ID	Description
1	37	100.0	6	ARI60918 Sequence
2	37	100.0	60	ARI60919 Sequence
3	37	100.0	6	ARI60918 Sequence
4	19	51.4	3	Continuation (4 of
5	19	51.4	5	Continuation (6 of
6	19	51.4	2	ACI02189 Mus muscu
7	19	51.4	2	ACI02189 Mus muscu
8	18	48.6	9	AL691449 Human DNA
9	18	48.6	2	AP001503 Homo sapi
10	18	48.6	9	AP000555 Homo sapi
11	18	48.6	2	AP002454 Mus muscu
12	18	48.6	2	AP002454 Homo sapi
13	18	48.6	2	AP002454 Homo sapi
14	18	48.6	2	AP001904 Homo sapi
15	18	48.6	2	ACI36510 Pan trogl
16	18	48.6	10	AL831776 Mouse DNA
17	18	48.6	9	ACI0928 Homo sapi
18	18	48.6	2	ACI09051 Rattus no
19	18	48.6	2	ACI15237 Rattus no

C 20	17	45.9	2997	9	AK127463	AK127463 Homo sapi
C 21	17	45.9	105856	9	AC015669	AC015669 Homo sapi
C 22	17	45.9	120817	2	ACI36057	ACI36057 Rattus no
C 23	17	45.9	149951	9	AC004970	AC004970 Homo sapi
C 24	17	45.9	156008	9	AL353093	AL353093 Human DNA
C 25	17	45.9	168602	2	ACI24099	ACI24099 Mus muscu
C 26	17	45.9	171112	9	AC016866	AC016866 Homo sapi
C 27	17	45.9	181936	2	AL360090	AL360090 Homo sapi
C 28	17	45.9	185472	2	ACI28825	ACI28825 Rattus no
C 29	17	45.9	201725	2	AL591706	AL591706 Homo sapi
C 30	17	45.9	206075	2	ACI18511	ACI18511 Rattus no
C 31	17	45.9	209380	10	ACI33186	ACI33186 Mus muscu
C 32	17	45.9	222162	10	ACI09281	ACI09281 Mus muscu
C 33	17	45.9	224226	2	ACI29414	ACI29414 Rattus no
C 34	17	45.9	235965	2	ACI22659	ACI22659 Rattus no
C 35	17	45.9	239924	2	ACI11266	ACI11266 Rattus no
C 36	17	45.9	240418	2	ACI28419	ACI28419 Rattus no
C 37	17	45.9	243598	2	ACI42076	ACI42076 Rattus no
C 38	17	45.9	243640	2	ACI08571	ACI08571 Rattus no
C 39	17	45.9	275059	2	ACI20776	ACI20776 Rattus no
C 40	16	43.2	666	11	BV047886	BV047886 Xenopus 1
C 41	16	43.2	1240	5	AF414086	AF414086 Xenopus 1
C 42	16	43.2	2420	5	CTD243835	CTD243835 Furobacte
C 43	16	43.2	12282	1	AE010462	AE010462 Furobacte
C 44	16	43.2	34667	5	BX571693	BX571693 Carp DNA
C 45	16	43.2	38490	9	AP001236	AP001236 Homo sapi

ALIGNMENTS

RESULT 1	ARI60918	Sequence 9 from patent US 6255281.	37 bp	DNA	linear	PAT 17-OCT-2001
LOCUS	ARI60918					
DEFINITION	ARI60918					
ACCESSION	ARI60918.1	GI:16225987				
VERSION						
KEYWORDS						
SOURCE	Unknown.					
ORGANISM	Unclassified.					
REFERENCE	1 (bases 1 to 37)					
AUTHORS	Pilon,A.L., Mukherjee,A.B. and Zhang,Z.					
TITLE	Use of recombinant human uteroglobin in treatment of inflammatory					
JOURNAL	and fibrotic conditions					
FEATURES	Patent: US 6255281-A 9 03-JUL-2001;					
source	Location/Qualifiers					
	1..37					
	/organism="unknown"					
	/mol_type="unassigned DNA"					

Query Match	100.0%	Score 37;	DB 6;	Length 37;
Best Local Similarity	100.0%	Pred. No. 9.7e-12;		
Matches	37;	Conservative	0;	Mismatches 0;
				Indels 0;
				Gaps 0;
Qy	1	ATGGAAGAATGCTCAGCTTACGCTGTGCACTAAG	37	
Db	1	ATGGAAGAATGCTCAGCTTACGCTGTGCACTAAG	37	
RESULT 2	ARI60919/c	Sequence 10 from patent US 6255281.	60 bp	DNA
LOCUS	ARI60919			
DEFINITION	ARI60919			
ACCESSION	ARI60919.1	GI:16225990		
VERSION				
KEYWORDS				
SOURCE	Unknown.			
ORGANISM	Unclassified.			
REFERENCE	1 (bases 1 to 60)			
AUTHORS	Pilon,A.L., Mukherjee,A.B. and Zhang,Z.			

TITLE	Use of recombinant human uteroglobin in treatment of inflammatory and fibrotic conditions
JOURNAL	Patent: US 6255281-A 10 03-UTL-2001;
FEATURES	location/Qualifiers
source	1. .60

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/mol_type="unassigned DNA"
ORIGIN

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Query Match	100.0%	Score 37;	DB 6;	length 60;
Best Local Similarity	100.0%;	Pred. No. 9.4e-12;		
Matches 37;	Conservative 0;	Mismatches 0;	Indels 0;	Gaps 0;

QY 1 ATGGAAGAATCGCTCAGTCTAGCCTGTGCACTAAG 3
Db 37 ATGGAAGAATCGCTCAGTCTAGCCTGTGCACTAAG 1

RESULT 3
AC102518 3/c
WPCOMMENT
Comments: null, data 4, comments 10000 1000000 10000000

Fragment Name	Begin	End
AC102518.0	1	110000
AC102518.1	100001	210000
AC102518.2	200001	310000
AC102518.3	300001	396850

Continuation of 4 of 4 of AC102518 from base 300001 (AC102518 Mus musculus chromosome 6 c

Query Match	51.4%;	Score 19;	DB 2;	Length 96850;
Best Local Similarity	100.0%;	Pred. No. 0.76;		
Matches 19;	Conservative 0;	Mismatches 0;	Indels 0;	Gaps 0;

QY	14	CTCAGTCTAGCCTGTGCAA	32
Db	22796	CTCAGTCTAGCCTGTGCAA	22778

RESULT 4
AC092450_5
WPCOMMENT

Fragment Name	Begin	End
AC092450_0	1	110000
AC092450_1	100001	210000
AC092450_2	200001	310000
AC092450_3	300001	410000
AC092450_4	400001	510000
AC092450_5	500001	610000
AC092450_6	600001	710000
AC092450_7	700001	727300

Continuation (6 of 8) of AC092450 from base 500001 (AC092450 Homo sapiens chromosome 12)

Query Match	51.4%;	Score 19;	DB 2;	Length 110000;
Best Local Similarity	100.0%;	Pred. No. 0.75;		
Matches 19;	Conservative 0;	Mismatches 0;	Indels 0;	Gaps 0;

OY	14	CTCAGTCTAGCCGTGCAA	32
Db	17216	CTCAGTCTAGCCGTGCAA	17234

RESULT 5	AC102189/c	LOCUS	AC102189	143223 bp	DNA	linear	HTG 16-JUL-2003
DEFINITION		Mus musculus clone RF24-21c77, WORKING DRAFT SEQUENCE, 12 unordered pieces.					

ACCESSION	AC102189
VERSION	AC102189.3
KEYWORDS	HTG; HTGS_PHASE1; HTGS_DRAFT; HTGS_FULLTROP.
SOURCE	Mus musculus (house mouse)
ORGANISM	Mus musculus
	Eukaryota; Metazoa; Chordata; Craniata; Vertebrata; Euteleostomi;

REFERENCE	AUTHORS	JOURNAL	REFERENCE	AUTHORS
1 (bases 1 to 143233)	Birren, B., Nusbaum, C. and Lander, E.	Mus musculus, Clone RP24-216U7	2 (bases 1 to 143233)	Birren, B., Linton, L., Nusbaum, C., Lander, E., Ali, A., Allen, N., Anderson, S., Baya, N., Beckten, V., Boncompagni, P., Boulanger, M.,
Unpublished				

Burns, S. J., Bahna, J. N., Casselberry, V., Dogherty, V. L., Deacon, G., Galletti, D. M., Brown, A., Canatella, J., Campolongo, A., Chang, Y., Chantaro, B., Chespe, J., Colangelo, M., Collins, S., Collymore, A., Cook, A., Cooke, P., D'Amelio, K., Devar, K., Diaz, J. S., Dodge, S., Fero, S., Ferreira, P., Fitzhugh, W., Gage, D., Galagan, Y., Gargyala, S., Girde, S., Gord, S., Goyette, M., Graham, L., Grand, Pierre, N., Hagos, B., Haeftord, A., Horton, L., Hulme, W., Iliev, I., Johnson, R., Jones, C., Kamat, A., Karatas, A., Kells, C., Lacroque, K., Lamazares, R., Landers, T., Lehoczy, J., Levine, R., Liu, G., Maclean, C., Macdonald, P., Major, J., Marcus, N., Matthews, C., McCarthy, M., Mcowan, P., McKernan, K., McSheeters, R., Meldrum, J., Menus, L., Mihova, T., Mianga, V., Murphy, T., Naylor, J., Nguyen, C., Norbu, C., Norman, C. H., O'Connor, T., O'Donnell, P., O'Neill, D., Oliver, J., Peterson, K., Phunkhang, P., Pierre, N., Pollara, V., Raymond, C., Retta, R., Rieback, M., Riley, R., Rise, C., Rogov, P., Roman, J., Rosettli, M., Roy, A., Santos, R., Schauer, S., Schnappack, R., Seman, S., Severly, P., Spencer, B., Stange-Thomann, N., Stojanovic, N., Strauss, N., Subramanian, A., Talamas, J., Teefaye, S., Theodore, J., Topfman, K., Travers, M., Trivis, N., Triggilo, J., Vassiliev, H., Vile, R., Vo, A., Wilson, B., Wu, X., Wyman, D., Ye, W. J., Young, G., Zainour, J., Zember, L., Zimmer, A., and Zody, M.

DIRECT SUBMISSION
 TITLE
 JOURNAL
 Submitted (23-NOV-2001) Whitehead Institute/MIT Center for Genome
 Research, 320 Charles Street, Cambridge, MA 02141, USA
 3 (bases 1 to 143233)
 REFERENCE
 Birren, B., Nusbaum, C., Lander, E., Abouelella, A., Allen, N.,
 AUTHORS

Anderson, M., Archchun, H. M., Hanna, N., Easten, V., Bloom, T., Boguslavsky, L., Boukgalter, B., Camerata, V., Chang, J., Choedel, Y., Collimore, A., Cook, A., Cooke, P., Corum, B., D'arellano, K., Diaz, J. S., Dodges, S., Doolley, K., Dorris, L., Erickson, J., Faro, S., Ferreira, P., Fitzgerald, M., Gage, D., Galagan, J., Garkyna, S., Gerham, L., Grand-Pierre, N., Hafez, N., Hasopian, D., Haos, B., Hall, J., Horton, L., Hulme, M., Iliev, I., Johnson, R., Jones, C., Kamat, A., Karatas, A., Kells, C., Landers, T., Levine, R., Limblad-Poh, K., Liu, X., Lui, A., Mabbitt, R., Maclean, C., Macdonald, P., Majer, J., Manning, J., Matthews, C., McCarthy, M., McElm, J., Menus, L., Mihova, T., Mierga, V., Murphy, T., Naylor, J., Nguyen, C., Nicol, R., Nobu, C., O'Connor, T., O'Donnell, P., O'Neill, D., Oliver, J., Peterson, K., Phunkhan, P., Pierre, N., Ratchupa, A., Ramasamy, U., Raymond, C., Retta, R., Risse, C., Rogov, P., Roman, J., Schauer, S., Schuppach, R., Seaman, S., Severy, P., Smith, C., Spencer, B., Stange-Thomann, N., Stojanovic, N., Stubbs, M., Talmes, J., Teatye, S., Theodore, J., Topham, K., Tvers, M., Vassiliou, H., Venkatacham, V. S., Viel, R., Vo, A., Wilson, B., Wu, X., Wyman, D., Young, G., Zainoun, J., Zembek, L., Zimmer, A. and Zody, M.

COMMENT JOURNAL
Submitted (16-JUN-2003) Whitehead Institute/MIT center for Genome
Research, 320 Charles Street, Cambridge, MA 02141 USA
On Jul 16, 2003 this sequence version replaced gi:22381210.

Smit, A.F.A. & Green, P. (1996-1997)
<http://ftp.genome.washington.edu/RM/RepeatMasker.html>
 Genome Center

Center: Whitehead Institute/ MIT Center for Genome Research
Center code: W1BR
Web site: <http://www.seq.wi.mit.edu>
Contact: sequence_submissions@genome.wi.mit.edu
----- Project Information -----
Center project name: L18187
Center clone name: 216_U7

```
----- Summary Statistics -----
Sequencing vector: plasmid, n/a; 100% of reads
Chemistry: Dye-terminator Big Dye; 100% of reads
Assembly program: Phrap; version 0.960731
Consensus quality: 139728 bases at least Q40
Consensus quality: 14119 bases at least Q30
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Consensus quality: 141907 bases at least Q20
Insert size: 133000; agarose-fp
Insert size: 142133; sum-of-contigs
Quality coverage: 10.7 in Q20 bases; agarose-fp
Quality coverage: 10.0 in Q20 bases; sum-of-contigs

```
* NOTE: This is a 'working draft' sequence. It currently
* consists of 12 contigs. The time order of the pieces
* is not known and their order in this sequence record is
* arbitrary. Gaps between the contigs are represented as
* runs of 'N', but the exact sizes of the gaps are unknown.
* This record will be updated with the finished sequence
* as soon as it is available and the accession number will
* be preserved. ....
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*	1	10233:	contig of 10232 bp in length
*	10233	gap of 100 bp	
*	10333	contig of 1103 bp in length	
*	11336	gap of 100 bp	
*	11536	contig of 12883 bp in length	
*	12334	gap of 100 bp	
*	12933	gap of 100 bp	
*	12934	contig of 2423 bp in length	
*	15357	gap of 100 bp	
*	15456	contig of 2210 bp in length	
*	15457	gap of 100 bp	
*	17667	contig of 2247 bp in length	
*	17667	gap of 100 bp	
*	20014	contig of 100 bp	
*	20114	contig of 3438 bp in length	
*	23852	gap of 100 bp	
*	23852	contig of 4654 bp in length	
*	27816	gap of 100 bp	
*	27916	contig of 8528 bp in length	
*	36443	gap of 100 bp	
*	36444	contig of 10224 bp in length	
*	46757	gap of 100 bp	
*	46767	contig of 2023 bp in length	
*	48858	gap of 100 bp	
*	48891	contig of 76243 bp in length	
*	65991	gap of 100 bp	
*	65991	contig of 100 bp	

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/mol_type="genomic DNA"
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/clone="RP24-216J7"
/clone_id="RRC1-24 Male Mouse BAC
1. 10232
/note="assembly_fragment
clone_end:596
vector_id:left"
10333. 11435
/note="assembly_fragment"
11536. 12813
/note="assembly_fragment"
12934. 15356
/note="assembly_fragment"
15457. 17666
/note="assembly_fragment"
17767. 20013
/note="assembly_fragment"
20114. 23551
/note="assembly_fragment"
23652. 27813
/note="assembly_fragment"
27916. 36443
/note="assembly_fragment"
36544. 46767
/note="assembly_fragment"
46868. 66890
/note="assembly_fragment"
66991. 143293
/note="assembly_fragment"

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Query Match	51.4%; Score 19; DB 2; Length 143233
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	Best Local Similarity	100.0%	Pred. No.	0.74;
	Matches	19;	Conservative	0;
	Mismatches		Indels	0;
	Gaps			0;
QY	14	CTCAGTCTAGGCTGTGCAA	32	
Db	105062	CTCAGTCTAGGCTGTGCAA	105044	

RESULT 6
AC092825/c

LOCUS	157842 bp	DNA	linear	PRI 22-MAY-2002
DEFINITION	AC0092825	3 BAC P11-171G7 (Roswell Park Cancer Institute Human		
ACCESSION	AC0092825	complete sequence.		

VERSION	AC092825.5	GI:21070529
KEYWORDS	HTG.	
SOURCE	Homo sapiens (human)	
ORGANISM	Homo sapiens	

REFERENCE
AUTHORS

Eukaryotes; Metazoa; Chordata; Craniata; Vertebrata; Euteleostomi; Mammalia; Eutheria; Primates; Catarrhini; Hominae; Homo.
1 (bases 1 to 157842)

Muzny, D.M., Adams, C., Adio-Oduola, B., Ali-osman, F.R., Allen, C.,

TITLE	Direct Submission
JOURNAL	Unpublished
REFERENCE	2 (bases 1 to 157842)
AUTHORS	Worley, K.C.
TITLE	Direct Submission
JOURNAL	Submitted (30-JUN-2001)
REFERENCE	CF Molecular and Human Genetics, Baylor College of Medicine, One Baylor Plaza, Houston, TX 77030, USA
AUTHORS	3 (bases 1 to 157842)
TITLE	Worley, K.C.
REFERENCE	Direct Submission

JOURNAL

Submitted (09-MAY-2002) Human Genome Sequencing Center, Department of Molecular and Human Genetics, Baylor College of Medicine, One Baylor Plaza, Houston, TX 77030, USA
4 (bases 1 to 157842)

REFERENCE
AUTHORS
TITLE
JOURNAL

Submitted (22-MAY-2002) Human Genome Sequencing Center, Department of Molecular and Human Genetics, Baylor College of Medicine, One Baylor Plaza, Houston, TX 77030, USA
On May 22, 2002 this sequence version replaced gi:20279217.
INFORMATION: <http://www.hgsc.bcm.tmc.edu/> or email gc-help@bcm.tmc.edu

COMMENT

CLONE LENGTH: This sequence does not necessarily represent the entire insert of this clone. Overlapping regions of clones are only sequenced and submitted once, so the sequence for the remainder of the insert may be found in the record for the adjacent clones. Overlapping clones are noted at the beginning and end of the Features listing.

ANNOTATION OF FEATURES:

STS are identified using ePCR (Genome Res. 7:541-550) searches of a local database that includes entries from dbSTS, GDB, and local mapping efforts.

Repeats are identified using RepeatMasker (A. Smit and P. Green, unpublished) for Human and Mouse sequences. Genes and Region of sequence similarity are identified by BLAST (Nuc. Acids Res. 25:3389-3402) similarity (expect < 1e-34) to the EST and cDNA sequences. Genes demonstrate at least two exons flanked by consensus splice sites that maintained sequence continuity across the splice junctions. Sequences that are not identical matches are annotated as similar.

SEQUENCING READ COVERAGE: Sequencing is completed to a minimum standard of double strand coverage with a minimum of 2 clones and 2 reads with no ambiguities or 2 chemistries with a minimum of 2 clones and 3 reads with no ambiguities. If the sequence quality for a region does not meet this standard, it will be indicated in the annotation as Low Coverage.

QUALITY OF INDIVIDUAL BASES: This sequence meets stringent quality standards - estimated error rate less than 1 per 10,000 bases. Reports of lowest quality individual bases and measures of base quality are listed below. Description of the metrics can be found at URL: <http://gc.bcm.tmc.edu:8088/quality.info/genbank.annotation.html>.

FEATURES

QUALSTAT-REPORT.

Location/Qualifiers

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   /mol_type="genomic DNA"
   /db_xref="taxon:9606"
   /chromosome="3"
   /clone="RP11-171G7"
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   /rpt_family="AT-rich"
   complement(206..514)
   /rpt_family="AluY"
   581..811
   /rpt_family="Charlie2"
   1005..1307
   /rpt_family="MLT2E"
   complement(1308..1371)
   /rpt_family="MER109"
   2881..3030
   /standard_name="81156"
   2824..2892
   /rpt_family="(TCTA)n"
   2899..2924
   /rpt_family="(CA)n"
   complement(3582..3979)
   /rpt_family="MSTRB"
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   complement(7020..7227)
   /rpt_family="MER101"
repeat_region 10026..10204
   /rpt_family="(TA)n"
   11515..11546
   /rpt_family="AT-rich"
   complement(11596..11904)
   /rpt_family="AluX"
   complement(15095..15165)
   /rpt_family="T199er1"
   15770..15789
   /rpt_family="(T)n"
   complement(15790..17993)
   /rpt_family="L1HS"
   17161..17338
   /standard_name="61535"
   18421..18480
   /rpt_family="L2"
   complement(21210..21359)
   /rpt_family="L1"
   23843..24385
   /rpt_family="MER1A"
   complement(24743..24873)
   /rpt_family="L1M3a"
   complement(25825..25997)
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   26400..26421
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   complement(26586..26814)
   /rpt_family="L2"
   27101..27277
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   complement(27621..28060)
   /rpt_family="L2"
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   complement(31014..31089)
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   complement(32739..32905)
   /rpt_family="MTR11"
   33642..33933
   /rpt_family="AluSg"
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   /rpt_family="AluX"
   35570..35756
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   37096..37118
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Query Match 51.4%; Score 19; DB 9; Length 157842;
Best Local Similarity 100.0%; Pred.No. 0.73;
Matches 19; Conservative 0; Mismatches 0; Indels 0; Gaps 0;
QY 14 CTCAGCTAGCGCTGCA 32
DB 117040 CTCAGCTAGCGCTGCA 117022

RESULT 7
 LOCUS AL691449 163934 bp DNA linear PRI 21-JAN-2003
 DEFINITION Human DNA sequence from clone RP11-558F24 on chromosome 1, complete sequence.
 ACCESSION AL691449
 VERSION AL691449.21 GI:27817297
 KEYWORDS HTG
 SOURCE Homo sapiens (human)
 ORGANISM Homo sapiens
 Eukaryota; Metazoa; Chordata; Craniata; Vertebrata; Euteleostomi; Mammalia; Eutheria; Primates; Catarrhini; Homnidae; Homo.
 REFERENCE 1 (bases 1 to 163934)
 AUTHORS Matthews, N.
 TITLE Direct Submission
 JOURNAL Submitted (21-JAN-2003) Wellcome Trust Sanger Institute, Hinxton, Cambridgeshire, CB10 1SA, UK. E-mail enquiries: humquerry@sanger.ac.uk
 On Jan 21, 2003 this sequence version replaced gi:27803162.
 COMMENT
 ----- Genome Center
 Center: Wellcome Trust Sanger Institute
 Center code: SC
 Web site: <http://www.sanger.ac.uk>
 Contact: humquerry@sanger.ac.uk

During sequence assembly data is compared from overlapping clones. Where differences are found these are annotated as variations together with a note of the overlapping clone name. Note that the variation annotation may not be found in the sequence submission corresponding to the overlapping clone, as we submit sequences with only a small overlap as described above.
 This sequence was finished as follows unless otherwise noted: all regions were either double-stranded or sequenced with an alternate chemistry or covered by high quality data (i.e., phred quality > 30); an attempt was made to resolve all sequencing problems, such as compressions and repeats; all regions were covered by at least one plasmid subclone or more than one M13 subclone; and the assembly was confirmed by restriction digest, except on the rare occasion of the clone being a YAC.
 The following abbreviations are used to associate primary accession numbers given in the feature table with their source databases: Em, EMBL; Sw, SWISSPROT; Tr, TREMBL; Wp, WORMPEP; Information on the WORMPEP database can be found at http://www.sanger.ac.uk/Projects/C_elegans/wormpep This sequence was generated from part of bacterial clone contigs of human chromosome 1, constructed by the Sanger Centre Chromosome 1 Mapping Group. Further information can be found at <http://www.sanger.ac.uk/HGP/Chr1>
 RP11-558F24 is from the library RP11-11.2 constructed by the group of Pletier de Jong. For further details see <http://www.chori.org/bacpac/home.htm>
 VECTOR: pBACe3.6.

FEATURES
 source location/Qualifiers
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 /mol_type="genomic DNA"
 /db_xref="taxon:9606"
 /chromosome="1"
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 /clone_11b="RP11-11.2"

Query Match 48.6%; Score 18; DB 9; Length 163934;
 Best Local Similarity 100.0%; Pred. No. 3;
 Matches 18; Conservative 0; Mismatches 0; Indels 0; Gaps 0;

QY 16 CAGCTAGCCTGTGCAC 33
 |||||
 DB 44909 CAGTCTAGCCTGTGCAC 44926

RESULT 8

AP001903
 LOCUS AP001903 167082 bp DNA linear HTG 30-MAY-2000
 DEFINITION Homo sapiens chromosome 18 clone RP11-719K4 map 18q21, WORKING DRAFT SEQUENCE, 26 unordered pieces.
 ACCESSION AP001903
 VERSION AP001903.2 GI:8117554
 KEYWORDS HTG; HTGS PHASE1; HTGS_DRAFT.
 SOURCE Homo sapiens (human)
 ORGANISM Homo sapiens
 Eukaryota; Metazoa; Chordata; Craniata; Vertebrata; Euteleostomi; Mammalia; Eutheria; Primates; Catarrhini; Homnidae; Homo.
 REFERENCE 1 (bases 1 to 167082)
 AUTHORS Hattori, M., Ishii, K., Toyoda, A., Taylor, T. D., Hong-Seog, P., Fujiyama, A., Yada, T., Totoki, Y., Watanabe, H. and Sakaki, Y.
 TITLE Published Only in Database (2000)
 JOURNAL 2 (bases 1 to 167082)
 Hattori, M., Ishii, K., Toyoda, A., Taylor, T. D., Hong-Seog, P., Fujiyama, A., Yada, T., Totoki, Y., Watanabe, H. and Sakaki, Y.
 Direct Submission
 Submitted (24-APR-2000) Masahira Hattori, The Institute of Physical and Chemical Research (RIKEN), Genomic Sciences Center (GSC), Kitasato Univ., 1-15-1 Kitasato, Sagami-hara, Kanagawa 228-8555, Japan (E-mail: hattori@gsc.riken.go.jp, URL: <http://hgp.gsc.riken.go.jp/>, Tel: 81-42-778-9923, Fax: 81-42-778-9924)
 On May 30, 2000 this sequence version replaced gi:7649790.
 COMMENT
 ----- Genome Center
 Center: RIKEN Genomic Sciences Center (GSC)
 Center code: RIKEN
 Web site: <http://hgp.gsc.riken.go.jp/>
 Contact: hattori@gsc.riken.go.jp
 ----- Project Information
 Center project name: HumDrat18
 Center clone name: RP11-719K4
 ----- Summary Statistics
 Sequencing vector: PCR products; 100% of reads
 Chemistry: Dye-terminator ET-amersham; 100% of reads
 Assembly program: Phrap; version 0.990329
 Consensus quality: 153234 bases at least Q40
 Consensus quality: 159520 bases at least Q30
 Consensus quality: 162786 bases at least Q20
 Insert size: 164582; sum-of-contrigs
 Quality coverage: 5.73x in Q20 bases; sum-of-contrigs

NOTE: This is a 'working draft' sequence. It currently consists of 26 contrigs. The true order of the pieces is not known and their order in this sequence record is arbitrary. Gaps between the contrigs are represented as runs N, but the exact sizes of the gaps are unknown. This record will be updated with the finished sequence as soon as it is available and the accession number will be preserved

1	22433 contrig of	22433 bp in length
22534	45017 contrig of	22443 bp in length
45118	59868 contrig of	14751 bp in length
59969	71793 contrig of	11825 bp in length
71894	81974 contrig of	10081 bp in length
82075	91440 contrig of	9366 bp in length
91541	98933 contrig of	6893 bp in length
98534	104713 contrig of	6180 bp in length
104814	108664 contrig of	3851 bp in length
108765	113626 contrig of	4862 bp in length
113727	120106 contrig of	6280 bp in length
120207	125902 contrig of	4766 bp in length
125103	129865 contrig of	4763 bp in length
129166	133326 contrig of	3361 bp in length
133427	138379 contrig of	4393 bp in length
138480	142887 contrig of	4408 bp in length
142988	146547 contrig of	3560 bp in length
146648	149630 contrig of	2863 bp in length
149631	152938 contrig of	3308 bp in length
153039	155111 contrig of	2073 bp in length
155212	157376 contrig of	2165 bp in length

157477 160277 contig of 2801 bp in length
 160378 162841 contig of 2464 bp in length
 162942 164597 contig of 1656 bp in length
 164698 166310 contig of 1613 bp in length
 166411 167082 contig of 672 bp in length

Sequence updated (26-May-2000).

* NOTE: This is a 'working draft' sequence. It currently consists of 26 contigs. The true order in this sequence record is not known and their order in this sequence record is arbitrary. Gaps between the contigs are represented as runs of N, but the exact sizes of the gaps are unknown. This record will be updated with the finished sequence as soon as it is available and the accession number will be preserved.

1 22433: contig of 22433 bp in length
 22434 22533: gap of 100 bp
 22534 45017: contig of 22484 bp in length
 45018 45117: gap of 100 bp
 45118 59668: contig of 14751 bp in length
 59669 59969: gap of 100 bp
 59970 71793: contig of 11825 bp in length
 71794 71893: gap of 100 bp
 71894 81974: contig of 10081 bp in length
 81975 91440: contig of 9366 bp in length
 91441 91540: gap of 100 bp
 91541 98433: contig of 6893 bp in length
 98434 98533: gap of 100 bp
 98534 104713: contig of 6180 bp in length
 104714 104813: gap of 100 bp
 104814 108664: contig of 3851 bp in length
 108665 108764: gap of 100 bp
 108765 113726: contig of 4862 bp in length
 113727 113727: gap of 100 bp
 113728 130106: contig of 6360 bp in length
 130107 120206: gap of 100 bp
 120207 125002: contig of 4756 bp in length
 125003 125102: gap of 100 bp
 125103 129865: contig of 4763 bp in length
 129866 129965: gap of 100 bp
 129966 133326: contig of 3361 bp in length
 133327 133426: gap of 100 bp
 133427 138379: contig of 4953 bp in length
 138380 138479: gap of 100 bp
 138480 142887: contig of 4408 bp in length
 142888 142987: gap of 100 bp
 142988 146547: contig of 3560 bp in length
 146548 146647: gap of 100 bp
 146648 149530: contig of 2883 bp in length
 149531 149630: gap of 100 bp
 149631 152938: contig of 3308 bp in length
 152939 153038: gap of 100 bp
 153039 153111: contig of 2073 bp in length
 153112 155211: gap of 100 bp
 155212 157376: contig of 2165 bp in length
 157377 157476: gap of 100 bp
 157477 160277: contig of 2801 bp in length
 160278 160377: gap of 100 bp
 160378 162841: contig of 2464 bp in length
 162842 162942: gap of 100 bp
 162943 164597: contig of 1656 bp in length
 164598 164698: gap of 100 bp
 164699 166310: contig of 1613 bp in length
 166311 166411: gap of 100 bp
 166412 167082: contig of 672 bp in length.

FEATURES
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1.167082
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 /mol_type="genomic DNA"
 /db_xref="taxon:9606"
 /chromosome="18"
 /map="18q21"
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 misc_feature 22534..45017
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 /note="assembly_fragment"
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 /note="assembly_fragment"
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 /note="assembly_fragment clone_end:5P6 vector_side:left"

ORIGIN

Query Match 48.6%; Score 18; DB 2; Length 167082;
 Best Local Similarity 100.0%; Pred. No. 3;
 Matches 18; Conservative 0; Mismatches 0; Indels 0; Gaps 0;

QY 16 CAGCTACGCTGTCGAC 33
 DB 70348 CAGCTACGCTGTCGAC 70365

RESULT 9 AP000555 168813 bp DNA linear PRI 01-OCT-1999
 LOCUS Homo sapiens genomic DNA, chromosome 22q11.2, BCL2 region,
 DEFINITION clone:RB1027C11.

ACCESSION AP000555
 VERSION AP000555.1 GI:5931541
 KEYWORDS
 SOURCE Homo sapiens (human)
 ORGANISM Homo sapiens

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Eukaryota; Metazoa; Chordata; Craniata; Vertebrata; Euteleostomi;
Mammalia; Eutheria; Primates; Catarrhini; Hominidae; Homo.
REFERENCE
1 (bases 1 to 168813)
AUTHORS
Shimizu,N.
TITLE
Human DNA sequence from clone KB1027C11 on chromosome 22q11.2
JOURNAL
Published Only in Database (1999)
REFERENCE
2 (bases 1 to 168813)
AUTHORS
Shimizu,N.
TITLE
Direct Submission
JOURNAL
Submitted (22-SEP-1999) Nobuyoshi Shimizu, Keio University, School
of Medicine, Molecular Biology, 35 Shinanomachi, Shinjuku-ku, Tokyo
160-0016, Japan (E-mail:nshimizu@med.keio.ac.jp,
Tel:81-3-3351-2370, Fax:81-3-3351-2370)
This is a complete sequence of the insert of KB1027C11 clone. The
proximal adjacent clone is KB665H9 (Acc.#AP000554) with 586-bp
overlapping. The distal adjacent clone is N109G12 (Acc.#D66995)
with 24893-bp overlapping.
FEATURES
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/cell_type="pre-pro-B cell"
/clone_lib="Keio BAC library"
3880..4003
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complement(6409..6701)
/evidence=not experimental
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complement(6814..6961)
/evidence=not experimental
/rpt_family="SFRM"
7486..7518
/evidence=not experimental
/rpt_family="AT-rich"
7839..8175
/evidence=not experimental
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20807..21027
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Best Local Similarity 100.0%; Pred. No. 3;
Matches 18; Conservative 0; Mismatches 0; Indels 0; Gaps 0;
Ox 16 CAGCTAGCTGTGCAAC 33
Db 73695 CAGCTAGCTGTGCAAC 73712
RESULT 10
AC102275 169230 bp DNA linear HTG 23-APR-2003
DEFINITION Mus musculus clone RP24-290E7, WORKING DRAFT SEQUENCE, 40 unordered
pieces.
AC102275
AC102275.2 GT30017900
KEYWORDS HTG; HTGS_PHASE1; HTGS_DRAFT.
SOURCE Mus musculus (house mouse)
ORGANISM Mus musculus
REFERENCE 1 (bases 1 to 169230)
AUTHORS Birren,B., Nusbaum,C. and Lander,E.
TITLE Mus musculus, clone RP24-290E7
JOURNAL Unpublished
REFERENCE 2 (bases 1 to 169230)

```

AUTHORS

Birren,B., Linton,L., Nusbaum,C., Lander,E., Ali,A., Allen,N., Anderson,S., Barna,N., Bastien,V., Boguslavsky,L., Bouhgalter,B., Brown,A., Camarata,J., Campopiano,A., Chang,J., Chararo,B., Chappel,V., Colangelo,M., Collins,S., Collymore,A., Cook,A., Cooke,P., Dearlano,K., Dewar,K., Diaz,J.S., Dodge,S., Fero,S., Ferreira,P., Fitzhugh,M., Gage,D., Galgan,J., Gardyna,S., Girde,S., Gora,S., Goyette,M., Graham,L., Grand-Pierre,N., Hagos,B., Heatford,A., Horton,L., Hulme,W., Iliev,I., Johnson,R., Jones,C., Kamat,A., Karatas,A., Kells,C., Lacroque,K., Lamazares,R., Landers,T., Lehoczy,J., Levine,N., Liu,G., Maclean,C., Macdonald,P., Major,J., Margus,N., Matthews,C., McCarthy,M., McEwan,P., McKernan,K., McPheters,R., Meldrum,C., Menus,L., Mohova,T., Mlenka,V., Murphy,T., Naylor,J., Nguyen,C., Norbu,C., Norman,C.H., O'Connor,T., O'Donnell,P., O'Neill,D., Oliver,J., Peterson,K., Phunkhang,P., Pierre,N., Pollara,V., Raymond,C., Retta,R., Rieback,W., Riley,R., Rise,C., Rogov,P., Roman,U., Rosetti,M., Roy,A., Santos,R., Schauer,S., Schupack,R., Seaman,S., Severy,P., Spencer,B., Stange-Thomann,N., Stojanovic,N., Strauss,N., Subramanian,A., Talamas,J., Tesfaye,S., Theodore,J., Topham,K., Travers,M., Travis,N., Triggillo,J., Vassiliev,H., Viel,R., Vo,A., Wilson,B., Wu,X., Wyman,D., Ye,W.J., Young,G., Zainoun,J., Zembek,L., Zimmer,A. and Zody,M.

TITLE

JOURNAL

REFERENCE

AUTHORS

Direct Submission
Submitted (23-NOV-2001) Whitehead Institute/MIT Center for Genome Research, 320 Charles Street, Cambridge, MA 02141, USA
3 (bases 1 to 169230)
Birren,B., Nusbaum,C., Lander,E., Abouelell,A., Allen,N., Anderson,S., Arachchi,H.M., Barna,N., Bastien,V., Bloom,T., Boguslavsky,L., Bouhgalter,B., Camarata,J., Chang,J., Chappel,Y., Collymore,A., Cook,A., Cooke,P., Corum,B., Dearlano,K., Diaz,J.S., Dodge,S., Dooley,K., Dorris,L., Erickson,J., Fero,S., Ferreira,P., Fitzhugh,M., Gage,D., Galgan,J., Gardyna,S., Graham,L., Grand-Pierre,N., Hagos,B., Iliev,I., Johnson,R., Jones,C., Hall,J., Horton,L., Hulme,W., Iliev,I., Johnson,R., Jones,C., Kamat,A., Karatas,A., Kells,C., Landers,T., Levine,R., Lindblad-Toh,K., Liu,G., Lui,A., Mabbitt,R., Maclean,C., Macdonald,P., Major,J., Manning,J., Matthews,C., McCarthy,M., Melrim,J., Menus,L., Mohova,T., Mlenka,V., Murphy,T., Naylor,J., Nguyen,C., Nicol,R., Nordu,C., O'Connor,T., O'Donnell,P., O'Neill,D., Oliver,J., Peterson,K., Phunkhang,P., Pierre,N., Rachupka,A., Ramasamy,U., Raymond,C., Retta,R., Rise,C., Rogov,P., Roman,U., Schauer,S., Schupack,R., Seaman,S., Severy,P., Smith,C., Spencer,B., Stange-Thomann,N., Stojanovic,N., Stubbs,M., Talamas,J., Tesfaye,S., Theodore,J., Topham,K., Travers,M., Vassiliev,H., Venkataraman,V.S., Viel,R., Vo,A., Wilson,B., Wu,X., Wyman,D., Young,G., Zainoun,J., Zembek,L., Zimmer,A. and Zody,M.
Direct Submission
Submitted (23-APR-2003) Whitehead Institute/MIT Center for Genome Research, 320 Charles Street, Cambridge, MA 02141, USA
On Apr 17, 2003 this sequence version replaced gi:17061361.
All repeats were identified using RepeatMasker:
Smit, A.F.A. & Green, P. (1996-1997)
<http://ftp.genome.washington.edu/RM/RepeatMasker.html>
----- Genome Center
Center: Whitehead Institute/ MIT Center for Genome Research
Center code: WIBR
Web site: <http://www-seq.wi.mit.edu>
Contact: sequence_submissions@genome.wi.mit.edu
----- Project Information
Center project name: L18289
Center clone name: 290_E_7

TITLE

JOURNAL

COMMENT

* NOTE: This is a 'working draft' sequence. It currently
* consists of 40 contigs. The true order of the pieces
* is not known and their order in this sequence record is
* arbitrary. Gaps between the contigs are represented as
* runs of N, but the exact sizes of the gaps are unknown.
* This record will be updated with the finished sequence
* as soon as it is available and the accession number will
* be preserved.
* 1 121664: contig of 121664 bp in length
* 121665 121764: gap of 100 bp
* 121765 122426: contig of 662 bp in length

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* 122427 122526: gap of 100 bp
* 122527 123632: contig of 1106 bp in length
* 123633 123732: gap of 100 bp
* 123733 124478: contig of 746 bp in length
* 124479 124578: gap of 100 bp
* 124579 125233: contig of 655 bp in length
* 125234 125233: gap of 100 bp
* 125334 126116: contig of 1083 bp in length
* 126117 126516: gap of 100 bp
* 126517 127253: contig of 737 bp in length
* 127254 128020: contig of 667 bp in length
* 128021 128120: gap of 100 bp
* 128121 128805: contig of 685 bp in length
* 128806 129774: gap of 100 bp
* 129775 129774: contig of 869 bp in length
* 129775 129874: gap of 100 bp
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* 130557 130556: gap of 100 bp
* 130557 131603: contig of 947 bp in length
* 131604 131703: gap of 100 bp
* 131704 132737: contig of 1034 bp in length
* 132738 132837: gap of 100 bp
* 132838 133498: contig of 661 bp in length
* 133499 133598: gap of 100 bp
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* 134341 134440: gap of 100 bp
* 134441 135439: contig of 899 bp in length
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* 136403 137116: contig of 714 bp in length
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* 137843 137942: gap of 100 bp
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* 139265 139364: gap of 100 bp
* 139365 140707: contig of 1343 bp in length
* 140708 140807: gap of 100 bp
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* 141957 142056: gap of 100 bp
* 142057 142855: contig of 799 bp in length
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* 150916 151015: gap of 100 bp
* 151016 152223: contig of 1208 bp in length
* 152224 152323: gap of 100 bp
* 152324 153902: contig of 1579 bp in length
* 153903 154002: gap of 100 bp
* 154003 155546: contig of 1544 bp in length
* 155447 155646: gap of 100 bp
* 155647 156697: contig of 1051 bp in length
* 156698 156797: gap of 100 bp
* 156798 158478: contig of 1681 bp in length
* 158479 158578: gap of 100 bp
* 158579 159753: contig of 1175 bp in length
* 159754 159853: gap of 100 bp
* 159854 161008: contig of 1155 bp in length
* 161009 161108: gap of 100 bp
* 161109 162235: contig of 1127 bp in length
* 162236 162335: gap of 100 bp

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FEATURES

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* 163345 163444: gap of 100 bp
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vector side:left"

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Best Local Similarity 100.0%; Pred. No. 3;
Matches 18; Conservative 0; Mismatches 0; Indels 0; Gaps 0;
QY 11 TCCTCAGCTCAGCCTGT 28
Db 65094 TCCTCAGCTCAGCCTGT 65111

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RESULT 11

AP002454

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LOCUS Homo sapiens chromosome 18 clone RP11-019K4 map 18q21, WORKING
DEFINITION DRAFT SEQUENCE, 29 unordered pieces.

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ACCESSION

AP002454

VERSION

AP002454.1

KEYWORDS

HTG; HTGS; PHASE1; HTGS_DRAFT.

SOURCE

Homo sapiens

ORGANISM

Homo sapiens

REFERENCE

1 (bases 1 to 172921)

AUTHORS

Hattori, M., Ishii, K., Toyoda, A., Taylor, T. D., Hong-Seog, P., Fujiyama, A., Yada, T., Torok, Y., Watanabe, H. and Sakaki, Y.

TITLE

Published Only in Database (2000)

JOURNAL

Hattori, M., Ishii, K., Toyoda, A., Taylor, T. D., Hong-Seog, P., Fujiyama, A., Yada, T., Torok, Y., Watanabe, H. and Sakaki, Y.

AUTHORS

Submitted (02-JUN-2000) Masahira Hattori, The Institute of Physical and Chemical Research (RIKEN), Genomic Sciences Center (GSC), Kitasato Univ., 1-15-1 Kitasato, Sagamihara, Kanagawa 228-8535,

misc_feature 88367.95348
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 DEFINITION Homo sapiens chromosome 5 clone CTD-2185023, complete sequence.
 ACCESSION AC091843
 VERSION AC091843.3 GI:18958638
 KEYWORDS HTG.
 SOURCE Homo sapiens (human)
 ORGANISM Eukaryota; Metazoa; Chordata; Craniata; Vertebrata; Euteleostomi;
 Mammalia; Eutheria; Primates; Catarrhini; Homnidae; Homo.
 REFERENCE 1 (bases 1 to 177056)
 AUTHORS DOE Joint Genome Institute and Stanford Human Genome Center.
 TITLE Direct Submission
 JOURNAL Unpublished
 REFERENCE 2 (bases 1 to 177056)
 AUTHORS DOE Joint Genome Institute.
 TITLE Direct Submission
 JOURNAL Submitted (09-JUN-2001) Production Sequencing Facility, DOE Joint
 Genome Institute, 2800 Mitchell Drive, Walnut Creek, CA 94598, USA
 3 (bases 1 to 177056)
 REFERENCE DOE Joint Genome Institute and Stanford Human Genome Center.
 TITLE Direct Submission
 JOURNAL Submitted (27-FEB-2002) DOE Joint Genome Institute, 2800 Mitchell
 Drive, Walnut Creek, CA 94598, USA
 On Feb 27, 2002 this sequence version replaced gi:15290412.
 COMMENT Draft Sequence Produced by DOE Joint Genome Institute
 www.jgi.doe.gov
 www.jgi.doe.gov
 Finishing Completed at Stanford Human Genome Center
 www-shgc.stanford.edu
 Quality: Phrap Quality >=40 100% of sequence;

FEATURES
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 location/Qualifiers
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ORIGIN
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 Best Local Similarity 100.0%; Pred. No. 3;
 Matches 18; Conservative 0; Mismatches 0; Indels 0; Gaps 0;

QY 16 CAGCTACCTGTGCAAC 33
 DB 14662 CAGCTACCTGTGCAAC 14645

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 DEFINITION Homo sapiens chromosome 18 clone RP11-720N18 map 18.
 ACCESSION AC090210
 VERSION AC090210.4 GI:15290848
 KEYWORDS HTG; HTGS PHASE2; HTGS_FULFILL; HTGS_CANCELLED.
 SOURCE Homo sapiens (human)
 ORGANISM Homo sapiens

REFERENCE 1 (bases 1 to 180851)
 AUTHORS Birren, B., Linton, L., Nusbaum, C. and Lander, E.
 TITLE Unpublished
 JOURNAL
 REFERENCE 2 (bases 1 to 180851)
 AUTHORS Birren, B., Linton, L., Nusbaum, C., Lander, E., Allen, N., Anderson, S.,

Barna, N., Biedt, V., Boguslavsky, L., Bouckgalter, B., Brown, A.,
 Camarata, J., Campopiano, A., Choepel, Y., Colangelo, M., Collins, S.,
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 Dodge, S., Faro, S., Ferreira, P., Fitzhugh, W., Gage, D., Galagan, J.,
 Gardyna, S., Ginde, S., Goyette, M., Graham, L., Grand-Pierre, N.,
 Hages, B., Heaford, A., Horton, L., Hulme, W., Iliev, I., Johnson, R.,
 Jones, C., Karatas, A., Labocque, K., Lamazares, R., Landers, T.,
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 Murphy, T., Naylor, J., Nguyen, C., Norbu, C., Norman, C., H.,
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 Wilson, B., Wu, X., Wyman, D., Ye, W., J., Young, G., Zainum, J.,
 Zembek, L., Zimmer, A. and Zody, M.
 TITLE Direct Submission
 JOURNAL Submitted (17-FEB-2001) Whitehead Institute/MIT Center for Genome
 Research, 320 Charles Street, Cambridge, MA 02141, USA
 REFERENCE 3 (bases 1 to 180851)
 AUTHORS Birren, B., Linton, L., Nusbaum, C., Lander, E., Allen, N.,
 Anderson, S., Barna, N., Biedt, V., Boguslavsky, L., Bouckgalter, B.,
 Brown, A., Camarata, J., Campopiano, A., Chang, J., Chazaro, B.,
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Norbu, C., Norman, C.H., O'Connor, T., O'Donnell, P., O'Neill, D.,
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 Raymond, C., Rella, R., Rieback, M., Riley, R., Rise, C., Rogov, P.,
 Roman, J., Rosetti, M., Roy, A., Santos, R., Schauer, S., Schupbach, R.,
 Saman, S., Severy, P., Spencer, B., Stange-Thomann, N., Stojanovic, N.,
 Strauss, N., Subramanian, A., Talamas, U., Testafaye, S., Theodore, T.,
 Topham, K., Travers, M., Travis, N., Trigg, J., Vassiliev, H.,
 Viel, R., Vo, A., Wilson, B., Wu, X., Wyman, D., Ye, W.J., Young, G.,
 Zainoun, J., Zembek, L., Zimmer, A., and Zody, M.
 Direct Submission
 Submitted (27-MAR-2003) Whitehead Institute/MIT Center for Genome
 Research, 320 Charles Street, Cambridge, MA 02141, USA
 On Aug 26, 2001 this sequence version replaced gi:14192991.
 All repeats were identified using RepeatMasker:
 Smit, A.F.A. & Green, P. (1996-1997)
<http://ftp.genome.washington.edu/MN/RepeatMasker.html>

Center: Whitehead Institute/ MIT Center for Genome Research
 Center code: WIBR
 Web site: <http://www-seq.wi.mit.edu>
 Contact: sequence_submissions@genome.wi.mit.edu

Project Information
 Center project name: L1223
 Center clone name: 720_N18

NOTE: This is a 'working draft' sequence. It currently
 consists of 1 contigs. Gaps between the contigs
 are represented as runs of N. The order of the pieces
 is believed to be correct as given, however the sizes
 of the gaps between them are based on estimates that have
 been provided by the submitter.
 This sequence will be replaced
 by the finished sequence as soon as it is available and
 the accession number will be preserved.
 1 180851: contig of 180851 bp in length.

FEATURES
 source
 1. 180851
 /organism="Homo sapiens"
 /mol_type="genomic DNA"
 /db_xref="taxon:9606"
 /chromosome="18"
 /map="18"
 /clone="RP11-720N18"
 /clone_1b="RP11-720N18 Human Male BAC"

ORIGIN
 Query Match 48.6% Score 18; DB 2; Length 180851;
 Best Local Similarity 100.0%; Pred. No. 3;
 Matches 18; Conservative 0; Mismatches 0; Indels 0; Gaps 0;

Oy 16 CAGTCTAGCCTGTGCAC 33
 Db 33216 CAGTCTAGCCTGTGCAC 33199

RESULT 14
 AP001904/c 181211 bp DNA linear HTG 30-MAY-2000
 LOCUS Homo sapiens chromosome 18 clone RP11-720N18 map 18q21, WORKING
 DEFINITION DRAFT SEQUENCE, 15 unordered pieces.

ACCESSION AP001904
 VERSION AP001904.2 GI:8117555
 KEYWORDS HTG; HTGS PHASE1; HTGS_DRAFT.
 SOURCE Homo sapiens (human)

ORGANISM Homo sapiens
 Eukaryota; Metazoa; Chordata; Craniata; Vertebrata; Euteleostomi;
 Mammalia; Eutheria; Primates; Catarrhini; Homiidae; Homo.

REFERENCE 1 (bases 1 to 181211)
 Hattori, M., Ishii, K., Toyoda, A., Taylor, T.D., Hong-Seog, P.,
 Fujiyama, A., Yada, T., Totoki, Y., Watanabe, H. and Sakaki, Y.
 Homo sapiens 181,211 genomic DNA of 18q21

TITLE JOURNAL
 Published only in Database (2000)
 2 (bases 1 to 181211)

AUTHORS Hattori, M., Ishii, K., Toyoda, A., Taylor, T.D., Hong-Seog, P.,
 Fujiyama, A., Yada, T., Totoki, Y., Watanabe, H. and Sakaki, Y.
 TITLE JOURNAL
 Submitted (24-APR-2000) Masahira Hattori, The Institute of Physical
 and Chemical Research (RIKEN), Genomic Sciences Center (GSC);
 Kitasato Univ., 1-15-1 Kitasato, Sagamihara, Kanagawa 228-8555,
 Japan (E-mail: hattori@gsc.riken.go.jp,
uri:http://hgp.gsc.riken.go.jp/, Tel:81-42-778-9923,
 Fax:81-42-778-9924)

COMMENT On May 30, 2000 this sequence version replaced gi:7649791.

Center: RIKEN Genomic Sciences Center (GSC)

Center code: RIKEN

Web site: <http://hgp.gsc.riken.go.jp/>

Contact: hattori@gsc.riken.go.jp

Project Information

Center project name: HumDRAFT18

Center clone name: RP11-720N18

Summary Statistics

Sequencing vector: PCR products; 100% of reads

Chemistry: Dye-terminator ET-amersham; 100% of reads

Assembly program: Phrap; version 0.990329

Consensus quality: 171968 bases at least Q40

Consensus quality: 176258 bases at least Q30

Consensus quality: 177945 bases at least Q20

Insert size: 179811; sum-of-contigs

Quality coverage: 5.94x in Q20 bases; sum-of-contigs

NOTE: This is a 'working draft' sequence. It currently consists of
 15 contigs. The true order of the pieces is not known and their
 order in this sequence record is arbitrary. Gaps between the
 contigs are represented as runs of N, but the exact sizes of the gaps
 are unknown. This record will be updated with the finished sequence
 as soon as it is available and the accession number will be
 preserved.

1 33300 contig of 33300 bp in length
 33401 54981 contig of 21581 bp in length
 55082 73468 contig of 18387 bp in length
 73569 94430 contig of 20862 bp in length
 94531 113614 contig of 19084 bp in length
 113715 123984 contig of 10270 bp in length
 124085 137815 contig of 13721 bp in length
 137916 147429 contig of 9514 bp in length
 147530 154898 contig of 7212 bp in length
 154939 162210 contig of 7269 bp in length
 162311 168580 contig of 6270 bp in length
 168681 171531 contig of 2851 bp in length
 171632 175227 contig of 3356 bp in length
 175328 178652 contig of 3325 bp in length
 178753 181211 contig of 2459 bp in length

Sequence updated (26-May-2000).

NOTE: This is a 'working draft' sequence. It currently
 consists of 15 contigs. The true order of the pieces
 is not known and their order in this sequence record is
 arbitrary. Gaps between the contigs are represented as
 runs of N, but the exact sizes of the gaps are unknown.
 This record will be updated with the finished sequence
 as soon as it is available and the accession number will
 be preserved.

1 33300: contig of 33300 bp in length
 33301 33400: gap of 100 bp
 33401 54981: contig of 21581 bp in length
 54982 55081: gap of 100 bp
 55082 73468: contig of 18387 bp in length
 73469 73568: gap of 100 bp
 73569 94430: contig of 20862 bp in length
 94431 94530: gap of 100 bp
 94531 113614: contig of 19084 bp in length
 113615 113714: gap of 100 bp
 113715 123984: contig of 10270 bp in length
 123985 124084: gap of 100 bp
 124085 137815: contig of 13721 bp in length
 137816 137915: gap of 100 bp


```

* 137916 147429: contig of 9514 bp in length
* 147430 147529: gap of 100 bp
* 147530 154898: contig of 7369 bp in length
* 154899 154998: gap of 100 bp
* 154999 162210: contig of 7212 bp in length
* 162211 162310: gap of 100 bp
* 162311 168580: contig of 6270 bp in length
* 168581 168680: gap of 100 bp
* 168681 171531: contig of 2851 bp in length
* 171532 171631: gap of 100 bp
* 171632 175227: contig of 3596 bp in length
* 175228 175327: gap of 100 bp
* 175328 178652: contig of 3325 bp in length
* 178653 181211: contig of 2459 bp in length.
* 178753

```

```

FEATURES
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1. 181211
/organism="Homo sapiens"
/mol_type="genomic DNA"
/db_xref="taxon:9606"
/chromosome="18"
/map="18q21"
/clone="RP11-720N18"
1. 333300
/misc_feature
/notes="assembly_fragment"
33401. 34981
/misc_feature
/notes="assembly_fragment"
55082. 73468
/misc_feature
/notes="assembly_fragment clone_end:SP6 vector_side:right"
73569. 94430
/misc_feature
/notes="assembly_fragment"
94531. 113614
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/notes="assembly_fragment"
113715. 123984
/misc_feature
/notes="assembly_fragment"
124085. 137815
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/notes="assembly_fragment"
137916. 147429
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/notes="assembly_fragment"
147530. 154898
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/notes="assembly_fragment clone_end:r7 vector_side:right"
154999. 162210
/misc_feature
/notes="assembly_fragment"
162311. 168580
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/notes="assembly_fragment"
168681. 171531
/misc_feature
/notes="assembly_fragment"
171632. 175227
/misc_feature
/notes="assembly_fragment"
175328. 178652
/misc_feature
/notes="assembly_fragment"
178753. 181211
/notes="assembly_fragment"

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ORIGIN

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Query Match 48.6%; Score 18; DB 2; Length 181211;
Best Local Similarity 100.0%; Pred. No. 3;
Matches 18; Conservative 0; Mismatches 0; Indels 0; Gaps 0;

```

```

QY 16 CAGCTTAGCCTGTGCAAC 33
DB 180151 CAGCTTAGCCTGTGCAAC 180134

```

```

RESULT 15
AC136510
LOCUS AC136510 186110 bp DNA linear HTG 18-MAR-2004
DEFINITION Pan troglodytes clone tp43-27n3, WORKING DRAFT SEQUENCE, 6 ordered
pieces.
ACCESSION AC136510
VERSION AC136510.17 GI:45544719
KEYWORDS HTG; HTGS PHASE2; HTGS DRAFT.
SOURCE Pan troglodytes (chimpanzee)

```

```

ORGANISM Pan troglodytes
REFERENCE Eukaryota; Metazoa; Chordata; Craniata; Vertebrata; Euteleostomi;
Mammalia; Eutheria; Primates; Catarrhini; Homnidae; Pan.
AUTHORS Lau,C.C.Y. and Roe,B.A.
TITLE 1 (bases 1 to 186110)
JOURNAL Pan troglodytes BAC clone tp43-27n3
REFERENCE Unpublished
AUTHORS 2 (bases 1 to 186110)
TITLE Lau,C.C.Y. and Roe,B.A.
JOURNAL Direct Submission
REFERENCE Submitted (05-NOV-2002) Department Of Chemistry And Biochemistry,
The University Of Oklahoma, 620 Parrington Oval, Room 208, Norman,
OK 73019, USA
REFERENCE 3 (bases 1 to 186110)
AUTHORS Lau,C.C.Y. and Roe,B.A.
JOURNAL Direct Submission
REFERENCE Submitted (18-MAR-2004) Department Of Chemistry And Biochemistry,
The University Of Oklahoma, 620 Parrington Oval, Room 208, Norman,
OK 73019, USA
COMMENT ----- Genome Center
Center: Department Of Chemistry And Biochemistry
The University Of Oklahoma
Center code:DOXNOR

```

```

* NOTE: This is a 'working draft' sequence. It currently
* consists of 6 contigs. Gaps between the contigs
* are represented as runs of N. The order of the pieces
* is believed to be correct as given, however the sizes
* of the gaps between them are based on estimates that have
* provided by the submittor.
* This sequence will be replaced
* by the finished sequence as soon as it is available and
* the accession number will be preserved.
1 76466: contig of 76466 bp in length
* 76467 76566: gap of unknown length
* 76567 82125: contig of 5559 bp in length
* 82126 82225: gap of unknown length
* 82226 92187: contig of 9962 bp in length
* 92188 92288: gap of unknown length
* 92289 94597: contig of 2310 bp in length
* 94598 94697: gap of unknown length
* 94698 112016: contig of 17319 bp in length
* 112017 112117: gap of unknown length
* 112118 186110: contig of 73994 bp in length.

```

FEATURES

```

source
1. 186110
/organism="Pan troglodytes"
/mol_type="genomic DNA"
/db_xref="taxon:9598"
/clone="tp43-27n3"
/clone_lib="RPC1 - 43 Male Chimpanzee BAC Library"

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ORIGIN

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Query Match 48.6%; Score 18; DB 2; Length 186110;
Best Local Similarity 100.0%; Pred. No. 3;
Matches 18; Conservative 0; Mismatches 0; Indels 0; Gaps 0;

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```

QY 16 CAGCTTAGCCTGTGCAAC 33
DB 36383 CAGCTTAGCCTGTGCAAC 36400

```

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Search completed: December 22, 2004, 23:36:46
Job time : 531.357 secs

```

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GenCore version 5.1.6
Copyright (c) 1993 - 2004 CompuGen Ltd.

OM nucleic - nucleic search, using sw model

Run on: December 22, 2004, 19:14:28 ; Search time 130.588 Seconds
(without alignments)
1487.336 Million cell updates/sec

Title: US-09-898-616A-4
Perfect score: 37
Sequence: 1 atgagaagatcgctcagctcagcctgctgcaactaag 37

Scoring table: OLIGO_NUC
Gapop 60.0 , Gapext 60.0

Searched: 4134866 seqs, 2624710521 residues

Word size : 0

Total number of hits satisfying chosen parameters: 8269772

Minimum DB seq length: 0
Maximum DB seq length: 2000000000

Post-processing: Listing first 45 summaries

Database : N Geneseq 23Sep04:*

1: geneseqn1980s:*\n2: geneseqn1990s:*\n3: geneseqn2000s:*\n4: geneseqn2001as:*\n5: geneseqn2001bs:*\n6: geneseqn2002as:*\n7: geneseqn2002bs:*\n8: geneseqn2003as:*\n9: geneseqn2003bs:*\n10: geneseqn2003cs:*\n11: geneseqn2003ds:*\n12: geneseqn2004s:*

Pred. No. is the number of results predicted by chance to have a score greater than or equal to the score of the result being printed, and is derived by analysis of the total score distribution.

SUMMARIES

Result No.	Score	Query Match	Length	ID	Description
1	37	100.0	37	9 ABZ58373	ABZ58373 Human ute
2	37	100.0	37	12 ADL27629	ADL27629 Recombina
3	37	100.0	60	9 ABZ58374	ABZ58374 Human ute
4	37	100.0	60	12 ADL27630	ADL27630 Recombina
5	16	43.2	698	4 AAK69658	AAK69658 Human imm
6	16	43.2	699	4 AAK69659	AAK69659 Human imm
7	16	43.2	699	4 AAK69660	AAK69660 Human imm
8	15	40.5	477	9 ACH14100	ACH14100 Human adu
9	15	40.5	621	9 AA192367	AA192367 Human pol
10	15	40.5	675	10 ADD35086	ADD35086 Mouse mit
11	15	40.5	1043	2 AA224882	AA224882 Human sec
12	15	40.5	1089	2 AAK37460	AAK37460 Human sec
13	15	40.5	1601	3 AA28524	AA28524 Human op1
14	15	40.5	1756	10 ADD44906	ADD44906 Rat gene
15	15	40.5	2289	3 AA28526	AA28526 Human op1
16	15	40.5	2290	3 AA28527	AA28527 Human op1
17	15	40.5	2348	3 AA28525	AA28525 Human op1
18	15	40.5	2408	3 AA28523	AA28523 Human op1
19	15	40.5	2409	4 AA17441	AA17441 Human cdn
20	15	40.5	2423	10 ADD14783	ADD14783 Human sec
21	15	40.5	3460	10 ADE56935	ADE56935 Rat gene

22	15	40.5	3715	8 ADA15021	ADA15021 Murine an
23	15	40.5	4065	4 AAD16913	AA16913 Human pro
24	15	40.5	8003	6 ABA93404	ABA93404 B. mori t
25	15	40.5	13660	4 AA199126	AA199126 Human exc
26	15	40.5	13660	4 AAK81002	AAK81002 Human imm
27	15	40.5	13660	4 AAK67907	AAK67907 Human imm
28	15	40.5	13660	5 AA163476	AA163476 Human kid
29	15	40.5	16122	4 AAS22908	AAS22908 DNA encod
30	15	40.5	16341	4 AAS2002	AAS2002 DNA encod
31	15	40.5	28001	12 AD136729	AD136729 Genomic D
32	15	40.5	28001	12 AD136730	AD136730 Genomic D
33	15	40.5	28001	12 ADM93169	ADM93169 Human KOX
34	15	40.5	28001	12 ADM93170	ADM93170 Human KOX
35	15	40.5	78025	8 ABQ77404	ABQ77404 Human SEL
36	15	40.5	153170	12 AD017382	AD017382 Human sof
37	15	40.5	183610	8 ACF62736	ACF62736 Cancer ba
38	15	40.5	183610	8 ADB20851	ADB20851 MRP1 base
39	15	40.5	183610	10 ADB87940	ADB87940 Human UGT
40	15	40.5	183610	10 ADB86923	ADB86923 Human MDR
41	15	40.5	183610	10 ADB82114	ADB82114 Human MDR
42	15	40.5	186591	8 ACF62750	ACF62750 Cancer ba
43	15	40.5	186591	8 ADB20869	ADB20869 MRP1 base
44	15	40.5	186591	10 ADB87958	ADB87958 Human UGT
45	15	40.5	186591	10 ADB96941	ADB96941 Human MDR

ALIGNMENTS

RESULT 1	ABZ58373
ID	ABZ58373 standard; DNA; 37 BP.
XX	ABZ58373;
XX	28-APR-2003 (first entry)
XX	Human uteroglobin synthetie gene oligonucleotide 4.
XX	Human; uteroglobin; respiratory distress; antiinflammatory; antifibrotic;
KW	antiinflammatory; antidiastmatic; nephrotoxic; antineumatic;
KW	antiarthritic; ss.
XX	
OS	Homo sapiens.
OS	Synthetic.
XX	
PN	WO2003003979-A2.
XX	
PD	16-JAN-2003.
XX	
PF	02-JUL-2002; 2002WO-US020836.
XX	
PR	02-JUL-2001; 2001US-00898616.
XX	
PA	(CLAR-) CLARAGEN INC.
XX	
PI	Pilon AL, Welch RE;
XX	
DR	WPI; 2003-221527/21.
XX	
PT	Bacterial expression system for producing recombinant human uteroglobin
PT	for treating inflammatory and fibrotic conditions, comprises a synthetic
PT	gene which codes for human uteroglobin.
XX	
PS	Claim 1; Page 33; 127pp; English.
XX	
CC	The present sequence is that of oligonucleotide 4, which was used in the
CC	construction of a synthetic gene for the production of human uteroglobin
CC	(hug) in bacteria. Oligonucleotides 1-4 (see ABZ58370-73) were used to
CC	assemble the coding strand and oligonucleotides 5-8 (see ABZ58374-77) the
CC	complementary strand. The gene was assembled by annealing and ligation of
CC	the oligonucleotides. Because mature native hug has glutamic acid at its
CC	N-terminus, an initiator methionine was added to the N-terminus, and

CC codon usage was optimised for expression in bacteria. In an example from
CC the invention, the synthetic gene was cloned into plasmid pCG12 (see
CC AB58378) and recombinant hUG (see ABP72259) was produced in *Escherichia*
CC coli strain CG12. The invention relates generally to the production of
CC recombinant hUG by bacterial expression, protein purification and scaled-
CC up production according to current good manufacturing practices. The
CC recombinant hUG is useful for the treatment of inflammatory and fibrotic
CC conditions, such as neonatal respiratory distress syndrome and
CC bronchopulmonary dysplasia. It may also be used to treat conditions
CC associated with elevated phospholipase A2 levels such as pancreatitis,
CC acute renal failure, rheumatoid arthritis and asthma.

SO Sequence 37 BP; 11 A; 8 C; 10 G; 8 T; 0 U; 0 Other;

Query Match 100.0%; Score 37; DB 9; Length 37;
Best Local Similarity 100.0%; Pred. No. 4.8e-11;
Matches 37; Conservative 0; Mismatches 0; Indels 0; Gaps 0;

QY 1 ATGAGAGAGATCGCTCAGTCTAGCTGTGCAACTAAG 37
DB 1 ATGAGAGAGATCGCTCAGTCTAGCTGTGCAACTAAG 37

RESULT 2
ADL27629
ID ADL27629 standard; DNA; 37 BP.
AC ADL27629;
XX
XX 20-MAY-2004 (first entry)
DE
XX Recombinant human uteroglobin, rhUG, coding oligonucleotide #4.
XX Human; ss; recombinant human uteroglobin, rhUG;
KM bacterial expression system; rhUG master cell bank;
KM rhUG research seed bank; anti-inflammatory; secretory phospholipase A 2;
KM fibronectin; respiratory distress; inflammation; fibrotic disease.
XX Homo sapiens.
OS Synthetic.
OS
XX US2003207795-A1.
PN
XX
XX 06-NOV-2003.
PD
XX
XX 02-JUL-2002; 2002US-00187498.
PF
XX
XX 28-MAY-1997; 97US-00864357.
PR
XX 02-JUL-2001; 2001US-00898616.
XX
XX (PILON) PILON A L.
PA (WELCH) WELCH R W.
XX
XX Pilon AL, Welch RW;
PI
XX WPI; 2004-051527/05.
DR
XX
XX Bacterial expression system for production of recombinant human
PT uteroglobin comprising synthetic gene or human cDNA sequence which codes
PT for human uteroglobin.
XX
XX
XX Claim 1; SEQ ID NO 4; 64pp; English.

CC cryovial and storing the cryovial at a temperature below -60 degrees C),
CC expressing rhUG (comprising providing a production seed cell bank culture
CC comprising an expression vector capable of expressing rhUG, inoculating a
CC broth medium with the production seed cell bank culture to form an
CC inoculum, incubating the bacterial culture formed in step (b) from the
CC inoculum, incubating the inoculum formed from the inoculum with the
CC step (c) to form a fermentation culture, adding an induction agent to
CC the fermentation culture to induce the expression of rhUG and harvesting
CC the above fermentation culture), purifying rhUG, determining the potency
CC of rhUG in a sample, measuring in vitro anti-inflammatory effect arising
CC from inhibition or blocking of secretory phospholipase A 2 enzymes by
CC rhUG, measuring in vitro binding of rhUG to fibronectin, determining the
CC purity of rhUG, and a pharmaceutical composition comprising a purified
CC rhUG and a carrier or diluent. The bacterial expression system is useful
CC for producing a rhUG research seed bank or a pharmaceutical grade rhUG
CC drug substance. rhUG is safe to administer to a patient in respiratory
CC distress. The rhUG is useful for treating inflammation and fibrotic
CC diseases. The present sequence is a coding strand oligonucleotide used to
CC construct the synthetic rhUG gene.

SO Sequence 37 BP; 11 A; 8 C; 10 G; 8 T; 0 U; 0 Other;

Query Match 100.0%; Score 37; DB 12; Length 37;
Best Local Similarity 100.0%; Pred. No. 4.8e-11;
Matches 37; Conservative 0; Mismatches 0; Indels 0; Gaps 0;

QY 1 ATGAGAGAGATCGCTCAGTCTAGCTGTGCAACTAAG 37
DB 1 ATGAGAGAGATCGCTCAGTCTAGCTGTGCAACTAAG 37

RESULT 3
ABZ58374/c
ID ABZ58374 standard; DNA; 60 BP.
AC ABZ58374;
XX
XX 28-APR-2003 (first entry)
DE
XX Human uteroglobin synthetic gene oligonucleotide 5.
XX Human; uteroglobin; respiratory distress; antiinflammatory; antifibrotic;
KM antiinflammatory; antiaesthetic; nephrotropic; antithematic;
KM antiaesthetic; ss.
XX Homo sapiens.
OS Synthetic.
OS
XX WO2003003979-A2.
PN
XX
XX 16-JAN-2003.
PD
XX
XX 02-JUL-2002; 2002MO-US020836.
PF
XX
XX 02-JUL-2001; 2001US-00898616.
PR
XX
XX (CLAR-) CLARAGEN INC.
PA
XX Pilon AL, Welch RE;
PI
XX WPI; 2003-221527/21.
DR
XX
XX Bacterial expression system for producing recombinant human uteroglobin
PT for treating inflammatory and fibrotic conditions, comprises a synthetic
PT gene which codes for human uteroglobin.
XX
XX Example 1; Page 33; 127pp; English.

CC The present sequence is that of oligonucleotide 5, which was used in the
CC construction of a synthetic gene for the production of human uteroglobin
CC (rhUG) in bacteria. Oligonucleotides 1-4 (see ABZ58370-73) were used to
CC assemble the coding strand and oligonucleotides 5-8 (see ABZ58374-77) the

complementary strand. The gene was assembled by annealing and ligation of the oligonucleotides. Because mature hUG has glutamic acid at its N-terminus, an initiator methionine was added to the N-terminus, and its codon usage was optimised for expression in bacteria. In an example from the invention, the synthetic gene was cloned into plasmid pCG12 (see ABZ86378) and recombinant hUG (see ABP72259) was produced in *Escherichia coli* strain C412. The invention relates generally to the production of recombinant hUG by bacterial expression, protein purification and scaled-up production according to current good manufacturing practices. The recombinant hUG is useful for the treatment of inflammatory and fibrotic conditions, such as neonatal respiratory distress syndrome and bronchopulmonary dysplasia. It may also be used to treat conditions associated with elevated phospholipase A2 levels such as pancreatitis, acute renal failure, rheumatoid arthritis and asthma.

Sequence 60 BP; 13 A; 14 C; 15 G; 18 T; 0 U; 0 Other;

Query Match 100.0%; Score 37; DB 9; Length 60;
Best Local Similarity 100.0%; Pred. No. 4.8e-11;
Matches 37; Conservative 0; Mismatches 0; Indels 0; Gaps 0;

1 ATGGAGAAGATCGCTCACTTACGCTGTGCACTTAAG 37
37 ATGGAGAAGATCGCTCACTTACGCTGTGCACTTAAG 1

RESULT 4
ADL27630/C
ID ADL27630 standard; DNA; 60 BP.
XX
XX ADL27630;
XX
XX 20-MAY-2004 (first entry)
XX
DE Recombinant human uteroglobin, rhUG, non-coding oligonucleotide #1.
XX
XX Human; ss; recombinant human uteroglobin; rhUG;
XX bacterial expression system; rhUG master cell bank;
XX rhUG research seed bank; anti-inflammatory; secretory phospholipase A 2;
XX fibronectin; respiratory distress; inflammation; fibrotic disease.
XX
XX Homo sapiens.
XX Synthetic.
XX US2003207795-A1.
XX
XX 06-NOV-2003.
XX
XX 02-JUL-2002; 2002US-00187498.
XX
XX 28-MAY-1997; 97US-00864357.
XX 02-JUL-2001; 2001US-00898616.
XX
XX (PILO/) PILON A L.
XX (WELC/) WELCH R W.
XX
XX Pilon AL, Welch RW;
XX
XX WPI; 2004-051527/05.
XX
XX Bacterial expression system for production of recombinant human
XX uteroglobin comprising synthetic gene or human cDNA sequence which codes
XX for human uteroglobin.
XX
XX Example 1; SEQ ID NO 5; 64bp; English.
XX
XX The invention relates to a bacterial expression system for the production
XX of recombinant human uteroglobin (rhUG), comprising a synthetic gene or
XX human cDNA sequence which codes for human UG, constructed from the
XX oligonucleotides appearing as ADL27626-ADL27629, and which further
XX comprises Met-Ala-Ala at the N-terminus of the sequence. Also included
XX are producing an rhUG master cell bank (comprising inoculating a suitable
XX incubating broth with an aliquot portion of a rhUG research seed bank to

form a bacterial culture, incubating the bacterial culture, adding a cryoprotective to the bacterial culture to form a cryopreserved solution, transferring a portion of the cryopreserved solution to a cryovial and storing the cryovial at a temperature below -60 degrees C), expressing rhUG (comprising providing a production seed cell bank culture comprising an expression vector capable of expressing rhUG, inoculating a broth medium with the production seed cell bank culture to form an inoculum, incubating the bacterial culture formed in step (b), inoculating a large scale fermenter with the inoculum formed from the step (c) to form a fermentation culture, incubating the fermentation culture within the large scale fermenter, adding an induction agent to the fermentation culture to induce the expression of rhUG and harvesting the above fermentation culture), purifying rhUG, determining the potency of rhUG in a sample, measuring in vitro anti-inflammatory effect arising from inhibition or blocking of secretory phospholipase A 2 enzymes by rhUG, measuring in vitro binding of rhUG to fibronectin, determining the purity of rhUG, and a pharmaceutical composition comprising a purified rhUG and a carrier or diluent. The bacterial expression system is useful for producing a rhUG research seed bank or a pharmaceutical grade rhUG drug substance. rhUG is safe to administer to a patient in respiratory distress. The rhUG is useful for treating inflammation and fibrotic diseases. The present sequence is a non-coding strand oligonucleotide used to construct the synthetic rhUG gene.

Sequence 60 BP; 13 A; 14 C; 15 G; 18 T; 0 U; 0 Other;

Query Match 100.0%; Score 37; DB 12; Length 60;
Best Local Similarity 100.0%; Pred. No. 4.8e-11;
Matches 37; Conservative 0; Mismatches 0; Indels 0; Gaps 0;

1 ATGGAGAAGATCGCTCACTTACGCTGTGCACTTAAG 37
37 ATGGAGAAGATCGCTCACTTACGCTGTGCACTTAAG 1

RESULT 5
AAK69658/C
ID AAK69658 standard; DNA; 698 BP.
XX
XX AAK69658;
XX
XX 06-NOV-2001 (first entry)
XX
XX Human immune/haematopoietic antigen genomic sequence SEQ ID NO:24470.
XX
XX Human; immune; haematopoietic; immune/haematopoietic antigen; cancer;
XX cytostatic; gene therapy; vaccine; metastasis; ds.
XX
XX Homo sapiens.
XX WO200157162-A2.
XX
XX 09-AUG-2001.
XX
XX 17-JAN-2001; 2001WO-US001354.
XX
XX 31-JAN-2000; 2000US-0179065P.
XX 04-FEB-2000; 2000US-0180628P.
XX 24-FEB-2000; 2000US-0184664P.
XX 02-MAR-2000; 2000US-0186350P.
XX 16-MAR-2000; 2000US-0189874P.
XX 17-MAR-2000; 2000US-0190076P.
XX 18-APR-2000; 2000US-0198123P.
XX 19-MAY-2000; 2000US-0205515P.
XX 07-JUN-2000; 2000US-0209467P.
XX 28-JUN-2000; 2000US-0215135P.
XX 30-JUN-2000; 2000US-0216647P.
XX 07-JUL-2000; 2000US-0216880P.
XX 11-JUL-2000; 2000US-0217487P.
XX 11-JUL-2000; 2000US-0217496P.
XX 14-JUL-2000; 2000US-0218290P.
XX 26-JUL-2000; 2000US-0220963P.

CC sequences from the present invention. AAK54942 to AAK54950 and AAK82169
CC represent sequences used in the exemplification of the present invention
SQ Sequence 698 BP; 270 A; 116 C; 107 G; 205 T; 0 U; 0 Other;

Query Match 43.2%; Score 16; DB 4; Length 698;
Best Local Similarity 100.0%; Pred. No. 30;
Matches 16; Conservative 0; Mismatches 0; Indels 0; Gaps 0;

QY 17 AGCTAGCCTGTGCA 32
Db 664 AGCTAGCCTGTGCA 649

RESULT 6
AAK69659/c
ID AAK69659 standard; DNA; 699 BP.

AC AAK69659;

DT 06-NOV-2001 (first entry)

DE Human immune/haematopoietic antigen genomic sequence SEQ ID NO:24471.

KW Human; immune; haematopoietic; immune/haematopoietic antigen; cancer;
cytostatic; gene therapy; vaccine; metastasis; ds.

OS Homo sapiens.

FN WO200157182-A2.

PD 09-AUG-2001.

PE 17-JAN-2001; 2001WO-US001354.

PR 31-JAN-2000; 2000US-0179065P.

PR 04-FEB-2000; 2000US-0180628P.

PR 24-FEB-2000; 2000US-0184684P.

PR 02-MAR-2000; 2000US-0186350P.

PR 16-MAR-2000; 2000US-0189874P.

PR 17-MAR-2000; 2000US-0190076P.

PR 18-APR-2000; 2000US-0198123P.

PR 19-MAY-2000; 2000US-0205515P.

PR 07-JUN-2000; 2000US-0209467P.

PR 28-JUN-2000; 2000US-0214888P.

PR 30-JUN-2000; 2000US-0215135P.

PR 07-JUL-2000; 2000US-0216880P.

PR 11-JUL-2000; 2000US-0217487P.

PR 14-JUL-2000; 2000US-0218290P.

PR 26-JUL-2000; 2000US-0220963P.

PR 26-JUL-2000; 2000US-0220963P.

PR 14-AUG-2000; 2000US-0224518P.

PR 14-AUG-2000; 2000US-0224518P.

PR 14-AUG-2000; 2000US-0225213P.

PR 14-AUG-2000; 2000US-0225213P.

PR 14-AUG-2000; 2000US-0225213P.

PR 14-AUG-2000; 2000US-0225213P.

PR 01-SEP-2000; 2000US-0229344P.

PR 01-SEP-2000; 2000US-0229345P.

PR 05-SEP-2000; 2000US-0229509P.

PR 05-SEP-2000; 2000US-0229513P.

PR 06-SEP-2000; 2000US-0230437P.

PR 06-SEP-2000; 2000US-0230438P.

PR 08-SEP-2000; 2000US-0231242P.

PR 08-SEP-2000; 2000US-0231243P.

PR 08-SEP-2000; 2000US-0231244P.

PR 08-SEP-2000; 2000US-0231413P.

PR 08-SEP-2000; 2000US-0231414P.

PR 08-SEP-2000; 2000US-0232080P.

PR 08-SEP-2000; 2000US-0232081P.

PR 12-SEP-2000; 2000US-0231958P.

PR 14-SEP-2000; 2000US-0232377P.

PR 14-SEP-2000; 2000US-0232398P.

PR 14-SEP-2000; 2000US-0232399P.

PR 14-SEP-2000; 2000US-0232400P.

PR 14-SEP-2000; 2000US-0233063P.

PR 14-SEP-2000; 2000US-0233064P.

PR 14-SEP-2000; 2000US-0233065P.

PR 21-SEP-2000; 2000US-0234223P.

PR 21-SEP-2000; 2000US-0234274P.

PR 25-SEP-2000; 2000US-0234937P.

PR 25-SEP-2000; 2000US-0234938P.

PR 26-SEP-2000; 2000US-0235484P.

PR 27-SEP-2000; 2000US-0235834P.

PR 29-SEP-2000; 2000US-0235836P.

PR 29-SEP-2000; 2000US-0236327P.

PR 29-SEP-2000; 2000US-0236377P.

PR 29-SEP-2000; 2000US-0236399P.

PR 29-SEP-2000; 2000US-0236399P.

PR 29-SEP-2000; 2000US-0236399P.

PR 29-SEP-2000; 2000US-0236399P.

PR 29-SEP-2000; 2000US-0236399P.

PR 29-SEP-2000; 2000US-0236399P.

PR 29-SEP-2000; 2000US-0236399P.

PR 29-SEP-2000; 2000US-0236399P.

PR 29-SEP-2000; 2000US-0236399P.

PR 29-SEP-2000; 2000US-0236399P.

PR 29-SEP-2000; 2000US-0236399P.

PR 29-SEP-2000; 2000US-0236399P.

PR 29-SEP-2000; 2000US-0236399P.


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PR 14-SEP-2000; 2000US-0233065P.
PR 21-SEP-2000; 2000US-0234222P.
PR 21-SEP-2000; 2000US-0234274P.
PR 25-SEP-2000; 2000US-0234979P.
PR 25-SEP-2000; 2000US-0234998P.
PR 26-SEP-2000; 2000US-0234984P.
PR 27-SEP-2000; 2000US-0235834P.
PR 27-SEP-2000; 2000US-0235836P.
PR 29-SEP-2000; 2000US-0236327P.
PR 29-SEP-2000; 2000US-0236368P.
PR 29-SEP-2000; 2000US-0236369P.
PR 29-SEP-2000; 2000US-0236370P.
PR 02-OCT-2000; 2000US-0236802P.
PR 02-OCT-2000; 2000US-0237037P.
PR 02-OCT-2000; 2000US-0237038P.
PR 02-OCT-2000; 2000US-0237039P.
PR 02-OCT-2000; 2000US-0237040P.
PR 13-OCT-2000; 2000US-0239335P.
PR 13-OCT-2000; 2000US-0239337P.
PR 20-OCT-2000; 2000US-0240960P.
PR 20-OCT-2000; 2000US-0241221P.
PR 20-OCT-2000; 2000US-0241785P.
PR 20-OCT-2000; 2000US-0241786P.
PR 20-OCT-2000; 2000US-0241787P.
PR 20-OCT-2000; 2000US-0241808P.
PR 20-OCT-2000; 2000US-0241809P.
PR 01-NOV-2000; 2000US-0244826P.
PR 01-NOV-2000; 2000US-0244617P.
PR 08-NOV-2000; 2000US-0244747P.
PR 08-NOV-2000; 2000US-0244750P.
PR 08-NOV-2000; 2000US-0246476P.
PR 08-NOV-2000; 2000US-0246477P.
PR 08-NOV-2000; 2000US-0246478P.
PR 08-NOV-2000; 2000US-0246523P.
PR 08-NOV-2000; 2000US-0246524P.
PR 08-NOV-2000; 2000US-0246525P.
PR 08-NOV-2000; 2000US-0246527P.
PR 08-NOV-2000; 2000US-0246528P.
PR 08-NOV-2000; 2000US-0246529P.
PR 08-NOV-2000; 2000US-0246609P.
PR 08-NOV-2000; 2000US-0246610P.
PR 08-NOV-2000; 2000US-0246611P.
PR 17-NOV-2000; 2000US-0248207P.
PR 17-NOV-2000; 2000US-0248208P.
PR 17-NOV-2000; 2000US-0248209P.
PR 17-NOV-2000; 2000US-0249210P.
PR 17-NOV-2000; 2000US-0249211P.
PR 17-NOV-2000; 2000US-0249212P.
PR 17-NOV-2000; 2000US-0249213P.
PR 17-NOV-2000; 2000US-0249214P.
PR 17-NOV-2000; 2000US-0249215P.
PR 17-NOV-2000; 2000US-0249216P.
PR 17-NOV-2000; 2000US-0249217P.
PR 17-NOV-2000; 2000US-0249218P.
PR 17-NOV-2000; 2000US-0249219P.
PR 17-NOV-2000; 2000US-0249244P.
PR 17-NOV-2000; 2000US-0249245P.
PR 17-NOV-2000; 2000US-0249264P.
PR 17-NOV-2000; 2000US-0249265P.
PR 17-NOV-2000; 2000US-0249297P.
PR 17-NOV-2000; 2000US-0249298P.
PR 17-NOV-2000; 2000US-0249300P.
PR 01-DEC-2000; 2000US-0250160P.
PR 01-DEC-2000; 2000US-0250391P.
PR 05-DEC-2000; 2000US-0251030P.
PR 05-DEC-2000; 2000US-0251988P.
PR 05-DEC-2000; 2000US-0256719P.
PR 06-DEC-2000; 2000US-0251479P.
PR 08-DEC-2000; 2000US-0251865P.
PR 08-DEC-2000; 2000US-0251868P.
PR 08-DEC-2000; 2000US-0251869P.

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PR 08-DEC-2000; 2000US-0251989P.
PR 08-DEC-2000; 2000US-0251990P.
PR 11-DEC-2000; 2000US-0254097P.
PR 05-JAN-2001; 2001US-0259678P.
XX
XX (HUMA-) HUMAN GENOME SCI INC.
XX
XX Rosen CA, Barash SC, Ruben SM;
PI WPI; 2001-483426/52.
DR
XX
XX Nucleic acids encoding human immune/hematopoietic antigen polypeptides,
PT useful for preventing, diagnosing and/or treating cancers and metastasis.
PT
XX
XX Disclosure; SEQ ID NO 24472; 3071bp + Sequence Listing; English.
XX
XX AAK54951 to AAK64702 encode the human immune/hematopoietic antigen (I)
CC amino acid sequences given in AAK62170 to AAK61921. (I) have cytostatic
CC activity, and can be used in gene therapy and vaccine production. (I)
CC proteins and polynucleotides may be used in the prevention, diagnosis and
CC treatment of diseases associated with inappropriate (I) expression. For
CC example, they may be used to treat disorders associated with decreased
CC expression by rectifying mutations or deletions in a patient's genome
CC that affect the activity of (I) by expressing inactive proteins or to
CC supplement the patient's own production of (I). Additionally, (I)
CC polynucleotides may be used to produce the secreted (I), by inserting the
CC nucleic acids into a host cell and culturing the cell to express the
CC protein. (I) proteins and polynucleotides may be used to prevent,
CC diagnose and treat immune/hematopoietic-related diseases, especially
CC cancer and cancer metastases of hematopoietic-derived cells. AAK64703
CC to AAK67694 represent human immune/hematopoietic antigen genomic
CC sequences from the present invention. AAK54942 to AAK54950 and AAK62169
CC represent sequences used in the exemplification of the present invention
CC
XX
SQ Sequence 699 BP; 270 A; 115 C; 107 G; 207 T; 0 U; 0 Other;
Query Match 43.2%; Score 16; DB 4; Length 699;
Best Local Similarity 100.0%; Pred. No. 30;
Matches 16; Conservative 0; Mismatches 0; Indels 0; Gaps 0;
QY 17 AGTCTAGCCTGTGCAA 32
DB 665 AGTCTAGCCTGTGCAA 650
RESULT 8
ACH14100
ID ACH14100 standard; cDNA; 477 BP.
XX
XX ACH14100;
AC
XX
XX 13-OCT-2003 (first entry)
DT
XX
XX Human adult brain cDNA #1312.
DE
XX
XX Human, ss; sequencing by hybridisation; SM; expressed sequence tag; EST;
KM genome mapping; biodiversity; genetic disorder.
XX
XX Homo sapiens.
OS
XX
XX US2003073623-A1.
PN
XX
XX 17-APR-2003.
PD
XX
XX 30-JUL-2001; 2001US-00918995.
PF
XX
XX 30-JUL-2001; 2001US-00918995.
PR
XX
XX 30-JUL-2001; 2001US-00918995.
XX
XX (DRMA/) DRMANAC R T.
PA (LABA/) LABAT I.
PA (STRAC/) STRACHE-CRAIN B.
PA (DICK/) DICKSON M C.
PA (JONE/) JONES L W.

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XX Drmanac RT, Labat I, Scache-Crain B, Dickson MC, Jones LW;
 PI WPI; 2003-615964/58.
 XX
 XX New polynucleotide sequences obtained from various cDNA libraries, useful
 PT as hybridization probes, as oligomers for PCR, for chromosome and gene
 PT mapping, in the recombinant production of protein, or in generating
 PT antisense DNA or RNA.
 PS Claim 1: SEQ ID NO 1312; 44pp; English.
 XX
 CC The invention relates to an isolated polynucleotide comprising any one of
 CC 38043 cDNA sequences, appearing as ACh12789-ACH5081, whose sequence was
 CC determined by the technique of SBH (sequencing by hybridisation). Also
 CC included is a purified polypeptide comprising a sequence corresponding to
 CC a reading frame of the novel polynucleotide. The nucleic acid sequences
 CC are useful in diagnostics as expressed sequence tags (EST) for
 CC identifying expressed genes or for physical mapping of the human genome,
 CC in forensics, in assessing biodiversity, or in identifying mutations
 CC responsible for genetic disorders and other traits. The nucleotide
 CC sequences are also useful as hybridisation probes, as oligomers for PCR,
 CC for chromosome and gene mapping, in the recombinant production of
 CC protein, or in generating antisense DNA or RNA. The purified polypeptide
 CC is useful for generating antibodies specific for it. The present sequence
 CC is one of the 38043 isolated cDNA/EST sequences. Note: The sequence data
 CC for this patent did not form part of the printed specification, but was
 CC obtained in electronic format directly from USPTO at
 CC seqdata.uspto.gov/sequence.html?DocID=20030073623
 CC
 XX Sequence 477 BP, 101 A; 116 C; 194 G; 52 T; 0 U; 14 Other;
 SQ
 Query Match 40.5%; Score 15; DB 9; Length 477;
 Best Local Similarity 100.0%; Pred. No. 1.1e+02;
 Matches 15; Conservative 0; Mismatches 0; Indels 0; Gaps 0;
 QY 2 TGGAGAGATCGCTC 16
 Db 305 TGGAGAGATCGCTC 319
 XX
 RESULT 9
 AAI92367
 ID AAI92367 standard; cDNA; 621 BP.
 XX
 AC AAI92367;
 XX
 DT 06-NOV-2001 (first entry)
 XX
 DE Human polynucleotide SEQ ID NO 12427.
 XX
 KW Human; cytokine; cell proliferation; cell differentiation; gene therapy;
 KW vaccine; peptide therapy; stem cell growth factor; haematopoiesis;
 KW tissue growth factor; immunomodulatory; cancer; leukaemia;
 KW nervous system disorders; arthritis; inflammation; ss.
 XX
 OS Homo sapiens.
 XX
 PA WO200164835-A2.
 XX
 PD 07-SEP-2001.
 XX
 PF 26-FEB-2001; 2001WO-US004927.
 XX
 PR 28-FEB-2000; 2000US-00515126.
 XX
 PR 18-MAY-2000; 2000US-00577409.
 XX
 PA (HYSE-) HYSEQ INC.
 XX
 PI Tang YT, Liu C, Drmanac RT;
 XX WPI; 2001-514838/56.
 DR P-PSDB; AAC12436.
 DR

XX
 PT Isolated nucleic acids and polypeptides, useful for preventing diagnosing
 PT and treating e.g. leukemia, inflammation and immune disorders.
 XX
 PS Claim 1: SEQ ID NO 12427; 1399pp + Sequence listing; English.
 XX
 CC The invention relates to human polynucleotides (AAI9941-AAI9841) and
 CC the encoded proteins (AAO0010-AAO13910) that exhibit activity elating to
 CC cytokine, cell proliferation or cell differentiation or which may induce
 CC production of other cytokines in other cell populations. The
 CC polynucleotides and polypeptides are useful in gene therapy, vaccines or
 CC peptide therapy. The polypeptides have various cytokine-like activities,
 CC e.g. stem cell growth factor activity, haematopoiesis regulating
 CC activity, tissue growth factor activity, immunomodulatory activity and
 CC activity/inhibin activity and may be useful in the diagnosis and/or
 CC treatment of cancer, leukaemia, nervous system disorders, arthritis and
 CC inflammation. Note: The sequence data for this patent did not form part
 CC of the printed specification, but was obtained in electronic format
 CC directly from WIPO at ftp.wipo.int/pub/published_pct_sequences
 XX
 SQ Sequence 621 BP, 152 A; 162 C; 137 G; 168 T; 0 U; 2 Other;
 XX
 Query Match 40.5%; Score 15; DB 4; Length 621;
 Best Local Similarity 100.0%; Pred. No. 1.1e+02;
 Matches 15; Conservative 0; Mismatches 0; Indels 0; Gaps 0;
 QY 19 TCTAGCCTGTGCAC 33
 Db 113 TCTAGCCTGTGCAC 127
 XX
 RESULT 10
 ADD35086
 ID ADD35086 standard; DNA; 675 BP.
 XX
 AC ADD35086;
 XX
 DT 15-JAN-2004 (first entry)
 XX
 DE Mouse mitochondrial DNA sequence SEQ ID NO:2866.
 XX
 KW ds; mouse; array; mitochondrial; hybridisation; energy metabolism;
 KW mitochondrial disease; oxidative phosphorylation dysfunction;
 KW oxidative stress; apoptosis; aging.
 XX
 OS Mus musculus.
 XX
 PA WO2003020220-A2.
 XX
 PD 13-MAR-2003.
 XX
 PF 30-AUG-2002; 2002WO-US027886.
 XX
 PR 30-AUG-2001; 2001US-0316323P.
 XX
 PR 31-AUG-2001; 2001CA-02356540.
 XX
 PA (UYEM-) UNIV EMORY.
 XX
 PI Wallace DC, Levy S, Kerstram K, Procaccio V;
 XX WPI; 2003-300821/29.
 XX
 PT Array containing probes for genes involved in mitochondrial biology,
 PT useful for determining mitochondrial biology gene expression profiles for
 PT use in diagnosing pathologies and identifying biochemical pathways.
 XX
 PS Claim 2: SEQ ID NO 2866; 201pp; English.
 XX
 CC The invention relates to a novel array comprising at least two isolated
 CC nucleotide molecules, each molecule having a sequence capable of uniquely
 CC hybridising to a nucleic acid molecule which is an expression product of
 CC a gene involved in mitochondrial biology. The array comprises two or more
 CC isolated nucleic acid molecules or spots, each molecule having a sequence

CC chosen from sequence of 994 human probes and 2046 mouse probes. An array
 CC of the invention is useful for determining an expression profile of a
 CC mouse or human sample containing nucleic acid by contacting the array
 CC with the sample under conditions allowing selective hybridization, and
 CC measuring hybridization of nucleic acid in the sample to the array to
 CC produce an expression profile. The array is also useful for determining
 CC an expression profile of a first labelled sample containing nucleic acid
 CC relative to a second, differently labelled sample containing nucleic
 CC acid. The second sample is a reference or a standard. An array is useful
 CC for determining an expression profile diagnostic of an energy-metabolism-
 CC related physiological condition. An array of the invention is useful for
 CC determining mitochondrial biology gene expression profiles of organisms,
 CC such as human, mice and closely related species, tissue and organs of
 CC such organisms, which are useful for determining expression profiles
 CC diagnostic of energy metabolism-related physiological conditions,
 CC diagnosing such physiological conditions, identifying biochemical
 CC pathways, genes, and mutations involved in such physiological conditions,
 CC identifying therapeutic agents useful for preventing and/or treating such
 CC physiological conditions, evaluating and/or monitoring the efficacy of
 CC such therapies, and creating and identifying animal models of human
 CC energy metabolism-related physiological conditions. An array is also
 CC useful for defining expression signatures or profiles for mitochondrial
 CC diseases, as well as distinguishing clinical disorders that result from
 CC oxidative phosphorylation (OXPHOS) dysfunction, oxidative stress,
 CC apoptosis and aging. An array of the invention contains probes of genes
 CC not previously recognised to participate in mitochondrial biology. The
 CC sequences shown in A033224-A033260 represent murine mitochondrial DNA
 CC clones used to make the probe of the invention. Some sequences are not
 CC present, these are SEQ ID NOs 295, 1174, 1213, 1700, 1728, 1730, 1905,
 CC 1906, 2408 and 2643.

CC Sequence 675 BP; 173 A; 142 C; 129 G; 221 T; 0 U; 0 Other;

CC Query Match 40.5%; Score 15; DB 10; Length 675;

CC Best Local Similarity 100.0%; Pred. No. 1.1e+02; Indels 0; Gaps 0;

CC Matches 15; Conservative 0; Mismatches 0; Indels 0; Gaps 0;

CC 15 TCAGCTCTAGCCTGTG 29

CC 301 TCAGCTCTAGCCTGTG 315

CC RESULT 11
 CC AA224882/C
 CC ID AA224882 standard; DNA; 1043 BP.

CC AA224882;

CC 02-DEC-1999 (first entry)

CC Human secreted protein gene 72 clone HB6GA29.

CC Human; secreted protein; fusion protein; gene therapy; protein therapy;
 CC diagnosis; tissue; cancer; tumour; neurodegenerative disorder; leukemia;
 CC developmental abnormality; foetal deficiency; blood; allergy; renal; ds;
 CC immune system; asthma; lymphocytic disease; brain; hepatic; lymphoma;
 CC inflammation; ischaemic shock; Alzheimer's disease; osteoarthritis; AIDS;
 CC cognitive disorder; schizophrenia; prostate; obesity; osteoarthritis; thymus;
 CC osteoporosis; arthritis; testis; lung; thyroiditis; thyroid digestion;
 CC endocrine; metabolism; regulation; malabsorption; gastritis; neoplasm.

CC Homo sapiens.

CC MO9947540-A1.

CC 23-SEP-1999.

CC 18-MAR-1999; 99WO-US005894.

CC 19-MAR-1999; 98US-0078563P.

CC 19-MAR-1999; 98US-0078563P.

CC 19-MAR-1999; 98US-0078573P.

CC 19-MAR-1999; 98US-0078574P.

PR 19-MAR-1998; 98US-0078576P.
 PR 19-MAR-1998; 98US-0078577P.
 PR 19-MAR-1998; 98US-0078578P.
 PR 19-MAR-1998; 98US-0078579P.
 PR 19-MAR-1998; 98US-0078581P.
 PR 01-APR-1998; 98US-0080312P.
 PR 01-APR-1998; 98US-0080313P.
 PR 01-APR-1998; 98US-0080314P.

PA (HUMA-) HUMAN GENOME SCI INC.

PI Ruben SM, Ni U, Rosen CA, Yu G, Young PE, Feng P, Soppet DR,

PI Wei Y, Endress GA, Duan RD, Kyaw H, Edner R, Lafleur DW, Olsen HS,

PI Shi Y, Moore PA.

DR WPI; 1999-562050/47.

DR F-PDSB; AAY41379.

PT New isolated human genes, useful for diagnosis and treatment of e.g.

PT cancers, neurological disorders, immune diseases, inflammation or blood

PT disorders.

PT Claim 1; Page 341; 484p; English.

CC This sequence represents a nucleic acid molecule which encodes a secreted
 CC human protein. The gene number, and the clone it is derived from, are
 CC detailed in the descriptor line. The gene can be used to generate fusion
 CC proteins by linking to the gene to a human immunoglobulin Fc portion
 CC (e.g. AA224882) for increasing the stability of the fused protein as
 CC compared to the human protein only. The invention relates to 95 novel
 CC genes and their fragments (nucleic acid sequences: AA224811-224907; amino
 CC acid sequences AAY41308-Y41404) which are useful for preventing, treating
 CC or ameliorating medical conditions e.g. by protein or gene therapy. Also,
 CC pathological conditions can be diagnosed by determining the amount of the
 CC new polypeptides in a sample or by determining the presence of mutations
 CC in the new polynucleotides. Specific uses are described for each of the
 CC 95 polynucleotides, based on which tissues they are most highly expressed
 CC in (see AA224811 for described uses)

CC Sequence 1043 BP; 281 A; 236 C; 291 G; 235 T; 0 U; 0 Other;

CC Query Match 40.5%; Score 15; DB 2; Length 1043;

CC Best Local Similarity 100.0%; Pred. No. 1.1e+02; Indels 0; Gaps 0;

CC Matches 15; Conservative 0; Mismatches 0; Indels 0; Gaps 0;

CC 19 TCTAGCCTGTGCAAC 33

CC 597 TCTAGCCTGTGCAAC 583

CC RESULT 12
 CC AA37460
 CC ID AA37460 standard; cDNA; 1089 BP.

CC AA37460;

CC 06-JUL-1999 (first entry)

CC Human secreted protein cDNA fragment containing gene 10.

CC Human; secreted protein; treatment; prevention; protein therapy; AIDS;
 CC gene therapy; diagnosis; cancer; tumour; neurodegenerative disorder;
 CC developmental abnormality; foetal deficiency; blood disorder; leukemia;
 CC immune system disease; autoimmune disease; hepatic disease; lymphoma;
 CC renal disease; inflammation; allergy; Alzheimer's disease; schizophrenia;
 CC cognitive disorder; prostate disease; skeletal; cardiac; muscle disorder;
 CC pulmonary disorder; transplant rejection; osteoarthritis; osteoporosis;
 CC arthritis; malignancy; digestive; endocrine; infection; ss.

CC Homo sapiens.

CC WO9918208-A1.

PD 15-APR-1999.
 XX
 PF 01-OCT-1998; 98MO-US020775.
 XX
 PR 02-OCT-1997; 97US-0060833P.
 PR 02-OCT-1997; 97US-0060836P.
 PR 02-OCT-1997; 97US-0060837P.
 PR 02-OCT-1997; 97US-0060838P.
 PR 02-OCT-1997; 97US-0060839P.
 PR 02-OCT-1997; 97US-0060843P.
 PR 02-OCT-1997; 97US-0060862P.
 PR 02-OCT-1997; 97US-0060866P.
 PR 02-OCT-1997; 97US-0060874P.
 PR 02-OCT-1997; 97US-0060880P.
 PR 02-OCT-1997; 97US-0060884P.
 PA (HUMA-) HUMAN GENOME SCI INC.
 XX
 PI Duan DR, Florence KA, Rosen CA, Ruben SM, Greene JM, Young P,
 PI Ferrie AM, Yu G, Janat F, Ni J, Carter KC, Endress GA, Feng P,
 PI Latleur DM, Shi Y;
 XX WPI; 1999-264022/22.
 DR P-PSDB; AAY07861.
 XX
 PT New isolated human genes and the secreted polypeptides they encode.
 XX
 PS Claim 1a; Page 232-233; 368pp; English.
 XX
 CC This invention describes novel isolated human genes and the secreted
 CC proteins they encode. The products of the invention are useful for
 CC preventing, treating or ameliorating medical conditions, e.g. by protein
 CC or gene therapy. Also pathological conditions can be diagnosed by
 CC determining the amount of the new polypeptides in a sample or by
 CC determining the presence of mutations in the new polynucleotides.
 CC Specific uses are described for each of the 101 polynucleotides, based on
 CC which tissues they are most highly expressed in, and include developing
 CC products for the diagnosis or treatment of cancer, tumours,
 CC neurodegenerative disorders, developmental abnormalities and fetal
 CC deficiencies, blood disorders, leukemias, diseases of the immune system,
 CC autoimmune diseases, hepatic and renal disease, lymphomas, inflammation,
 CC allergies, Alzheimer's and cognitive disorders, pulmonary disorders,
 CC disease, skeletal or cardiac muscle disorders, primary disorders,
 CC osteoporosis, arthritis or malignancies, digestive/endocrine disorders,
 CC infections and AIDS. The human secreted proteins of the invention are
 CC represented in AAY07852-Y07993 and the encoding nucleic acids are
 CC represented in AAX37451-X37552
 XX
 SO Sequence 1089 BP; 367 A; 202 C; 240 G; 279 T; 0 U; 1 Other;
 Query Match 40.5%; Score 15; DB 2; Length 1089;
 Best Local Similarity 100.0%; Pred. No. 1.1e+02;
 Matches 15; Conservative 0; Mismatches 0; Indels 0; Gaps 0;
 QY 3 GGAGAGATCGCTCA 17
 Db 14 GGAGAGATCGCTCA 28
 RESULT 13
 AAA28524
 ID AAA28524 standard; cDNA; 1601 BP.
 XX
 AC AAA28524;
 XX
 DT 29-AUG-2000 (first entry)
 XX
 DE Human opioid growth factor receptor cDNA spliced version 1.
 XX
 KW OGR; opioid growth factor receptor; growth inhibitor; proliferative;
 KW cytoskeletal; vulnery; gene therapy; antagonist; chromosome 20q13.3; ss.
 XX

OS Homo sapiens.
 FH Key Location/Qualifiers
 FT 1..33
 FT 5' UTR /tag= a
 FT CDS 34..1419
 FT /tag= b
 FT /product= "Opioid_growth_factor_receptor"
 FT 3' UTR 1420..1601
 FT /*tag= c
 EN WO00026340-A2.
 XX
 PD 11-MAY-2000.
 XX
 PF 02-NOV-1999; 99MO-US025802.
 XX
 PR 03-NOV-1998; 98US-0106879P.
 XX
 PA (PENN-) PENN STATE RES FOUND.
 XX
 PI Zagon IS, McLaughlin PJ, Verderame MF;
 XX
 XX WPI; 2000-365594/31.
 DR P-PSDB; AAY92810.
 XX
 PT New cDNA encoding rat and human opioid growth factor receptors which
 PT modulate cell growth, useful for treating cancer.
 XX
 PS Claim 1; Page 82-83; 91pp; English.
 XX
 CC Primers generated from rat opioid growth factor receptor (OGR) cDNA were
 CC used to clone a fragment of the human OGR cDNA. The complete sequence of
 CC human OGR was assembled with a combination of 3' and 5' RACE. 5' RACE
 CC consistently yielded a single species of cDNA, while the 3' RACE revealed
 CC extensive alternative splicing. The alternate splice forms were missing
 CC the imperfect repeats or differed in the number of imperfect repeats. The
 CC human OGR gene chromosomal location was determined by FISH as 20q13.3.
 CC OGR proteins, nucleic acid molecules, antibodies, transformed cells and
 CC expression vector are useful for detecting expression or levels of an
 CC OGR in a tissue. OGR nucleic acids can be used to inhibit growth of
 CC cells in vitro. The antisense sequences and antibodies can be used to
 CC promote growth of cells in vitro. Cell growth can be promoted by
 CC interfering with the OGR ligand-receptor system, especially where a
 CC subject suffers from a tissue wound. Treating cancer comprises enhancing
 CC the function of the OGR ligand-receptor system in cancerous cells of a
 CC patient or administering the OGR nucleic acid to the patient
 XX
 SO Sequence 1601 BP; 322 A; 485 C; 558 G; 236 T; 0 U; 0 Other;
 Query Match 40.5%; Score 15; DB 3; Length 1601;
 Best Local Similarity 100.0%; Pred. No. 1.1e+02;
 Matches 15; Conservative 0; Mismatches 0; Indels 0; Gaps 0;
 QY 2 TGGAGAGATCGCTC 16
 Db 1253 TGGAGAGATCGCTC 1267
 RESULT 14
 ADD44906/c
 ID ADD44906 standard; DNA; 1756 BP.
 XX
 AC ADD44906;
 XX
 DT 29-JAN-2004 (first entry)
 XX
 DE Rat gene Y15054, SEQ ID NO 10337.
 XX
 KW Rat; ds; gene; pain; neuronal tissue; gene therapy;
 KW spinal segmental nerve injury; chronic constriction injury; CCI;
 KW spared nerve injury; SNI; Chung.
 XX

OS Rattus norvegicus.
 XX
 XX WO2003016475-A2.
 XX
 XX 27-FEB-2003.
 XX
 PF 14-AUG-2002; 2002WO-US025765.
 XX
 XX 14-AUG-2001; 2001US-0312147P.
 PR 01-NOV-2001; 2001US-0346382P.
 PR 26-NOV-2001; 2001US-0333347P.
 XX
 PA (GENO) GEN HOSPITAL CORP.
 PA (FARO) BAYER AG.
 FI Woolf C, D'urso D, Befort K, Costigan M;
 XX WPI; 2003-268312/26.
 DR GENBANK; Y15054.
 XX
 PT New composition comprising two or more isolated polypeptides, useful for
 PT preparing a medicament for treating pain in an animal.
 XX
 PS Claim 1; Page: 1017P; English.

XX The invention discloses a composition comprising two or more isolated rat
 CC or human polynucleotides or a polynucleotide which represents a fragment,
 CC derivative or allelic variation of the nucleic acid sequence. Also
 CC claimed are a vector comprising the novel polynucleotide, a host cell
 CC comprising the vector, a method for identifying a nucleotide sequence
 CC which is differentially regulated in an animal subjected to pain and a
 CC kit to perform the method, an array, a method for identifying an agent
 CC that increases or decreases the expression of the polynucleotide sequence
 CC that is differentially expressed in neuronal tissue of a first animal
 CC subjected to pain, a method for identifying a compound which regulates
 CC the expression of a polynucleotide sequence which is differentially
 CC expressed in an animal subjected to pain, a method for identifying a
 CC compound that regulates the activity of one or more of the
 CC polynucleotides, a method for producing a pharmaceutical composition, a
 CC method for identifying a compound or small molecule that regulates the
 CC activity in an animal of one or more of the polypeptides given in the
 CC specification, a method for identifying a compound useful in treating
 CC pain and a pharmaceutical composition comprising the one or more
 CC polypeptides or their antibodies. The polynucleotide or the compound that
 CC modulates its activity is useful for preparing a medicament for treating
 CC pain (e.g. spinal segmental nerve injury (Chung), chronic constriction
 CC injury (CCI) and spared nerve injury (SNI)) in an animal (e.g. Gene
 CC therapy). The sequence presented is a rat DNA (shown in Table 2 of the
 CC specification) which encodes one of the polypeptides of the invention
 CC which is differentially expressed during pain. Note: The sequence data
 CC for this patent did not form part of the printed specification, but was
 CC obtained in electronic form directly from WIPO at
 CC ftp.wipo.int/pub/published_pct_sequences.

SO Sequence 1756 BP; 355 A; 543 C; 484 G; 374 T; 0 U; 0 Other;

Query Match 40.5%; Score 15; DB 10; Length 1756;

Best Local Similarity 100.0%; Pred. No. 1.1e+02;

Matches 15; Conservative 0; Mismatches 0; Indels 0; Gaps 0;

QY 13 GCTCAGCTAGCCTG 27
 DB 402 GCTCAGCTAGCCTG 388

RESULT 15
 AAA28526

ID AAA28526 standard; cDNA; 2289 BP.

XX AAA28526;

AC 29-AUG-2000 (first entry)

DT
 XX

DE Human opioid growth factor receptor cDNA spliced version 7.
 XX
 XX OGR, opioid growth factor receptor; growth inhibitor; proliferative;
 KM cytoskeletal; vulnertary, gene therapy; antagonist; chromosome 20q13.3; ss.
 XX
 OS Homo sapiens.
 XX
 FH Key Location/Qualifiers
 FT 5'UTR 1..33
 FT CDS /*tag= a
 FT 34..2007
 FT /*tag= b
 FT /*product= "Opioid_growth_factor_receptor"
 FT 2008..2289
 FT /*tag= c

WO200026340-A2.

11-MAY-2000.

02-NOV-1999; 99WO-US025802.

03-NOV-1998; 98US-0106879P.

(PENN-) PENN STATES RES FOUND.

Zagon IS, McLaughlin PJ, Verderame MF;

WPI; 2000-365594/31.

P-PDB; AA92812.

PT New cDNA encoding rat and human opioid growth factor receptors which
 PT modulate cell growth, useful for treating cancer.

PS Claim 1; Page 87-89; 91pp; English.

CC Primers generated from rat opioid growth factor receptor (OGR) cDNA were
 CC used to clone a fragment of the human OGR cDNA. The complete sequence of
 CC human OGR was assembled with a combination of 3' and 5' RACE. 5' RACE
 CC consistently yielded a single species of cDNA, while the 3' RACE revealed
 CC extensive alternative splicing. The alternate splice forms were missing
 CC the imperfect repeats or differed in the number of imperfect repeats. The
 CC human OGR gene chromosomal location was determined by FISH as 20q13.3.
 CC OGR proteins, nucleic acid molecules, antibodies, transformed cells and
 CC expression vector are useful for detecting expression or levels of an
 CC OGR in a tissue. OGR nucleic acids can be used to inhibit growth of
 CC cells in vitro. The antisense sequences and antibodies can be used to
 CC promote growth of cells in vitro. Cell growth can be promoted by
 CC interfering with the OGR ligand-receptor system, especially where a
 CC subject suffers from a tissue wound. Treating cancer comprises enhancing
 CC the function of the OGR ligand-receptor system in cancerous cells of a
 CC patient or administering the OGR nucleic acid to the patient

SO Sequence 2289 BP; 470 A; 714 C; 809 G; 296 T; 0 U; 0 Other;

Query Match 40.5%; Score 15; DB 3; Length 2289;

Best Local Similarity 100.0%; Pred. No. 1.1e+02;

Matches 15; Conservative 0; Mismatches 0; Indels 0; Gaps 0;

QY 2 TGGAGAGATGGCTC 16
 DB 1253 TGGAGAGATGGCTC 1267

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 Job time : 136.838 secs

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OM nucleic - nucleic search, using sw model

Run on: December 22, 2004, 22:08:48 ; Search time 29.8009 Seconds
(without alignments)
882.496 Million cell updates/sec

Title: US-09-898-616A-4

Perfect score: 37
Sequence: 1 atgagagaagatcgctcagctcgtcgaactaag 37

Scoring table: OLIGO_NUC
Gapop 60.0 , Gapext 60.0

Searched: 824507 seqs, 355394441 residues

Word size : 0

Total number of hits satisfying chosen parameters: 1649014

Minimum DB seq length: 0
Maximum DB seq length: 2000000000

Post-processing: Listing first 45 summaries

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5: /cgn2_6/ptodata/1/ina/PTCUS.COMB.seq.*
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Pred. No. is the number of results predicted by chance to have a score greater than or equal to the score of the result being printed, and is derived by analysis of the total score distribution.

SUMMARIES

Result No.	Score	Query Match	Length	DB ID	Description
1	37	100.0	37	US-08-864-357F-9	Sequence 9, Appli
2	37	100.0	60	US-08-864-357F-10	Sequence 10, Appli
3	15	40.5	3715	US-09-234-245-1	Sequence 1, Appli
4	14	37.8	283	US-08-760-534A-5	Sequence 5, Appli
5	14	37.8	283	US-09-336-757-5	Sequence 5, Appli
6	14	37.8	341	US-09-513-999C-3565	Sequence 3565, Ap
7	14	37.8	422	US-09-621-976-12461	Sequence 12461, A
8	14	37.8	553	US-08-463-115-26	Sequence 26, Appli
9	14	37.8	553	US-08-465-388-26	Sequence 26, Appli
10	14	37.8	715	US-08-463-115-18	Sequence 18, Appli
11	14	37.8	715	US-08-465-388-18	Sequence 18, Appli
12	14	37.8	1374	US-09-253-991A-5999	Sequence 5999, Ap
13	14	37.8	1540	US-08-463-115-2	Sequence 2, Appli
14	14	37.8	1540	US-08-465-388-2	Sequence 2, Appli
15	14	37.8	1899	US-09-253-991A-6306	Sequence 6306, Ap
16	14	37.8	2196	US-09-799-461-298	Sequence 298, App
17	14	37.8	2196	US-08-472-217-3	Sequence 3, Appli
18	14	37.8	2196	US-08-760-534A-3	Sequence 3, Appli
19	14	37.8	2196	US-09-336-757-3	Sequence 3, Appli
20	14	37.8	2895	US-09-556-877-171	Sequence 171, App
21	14	37.8	2895	US-09-620-412C-171	Sequence 171, App
22	14	37.8	2895	US-09-598-419-171	Sequence 171, App
23	14	37.8	2934	US-09-556-877-183	Sequence 183, App
24	14	37.8	2934	US-09-620-412C-183	Sequence 183, App
25	14	37.8	2934	US-09-598-419-183	Sequence 183, App
26	14	37.8	4000	US-09-780-049-18	Sequence 18, Appli
27	14	37.8	72604	US-09-268-992-7	Sequence 7, Appli

C 28	14	37.8	72604	3	US-09-657-474-7	Sequence 7, Appli
C 29	14	37.8	1830121	4	US-09-557-884-1	Sequence 1, Appli
C 30	14	37.8	1830121	4	US-09-643-990A-1	Sequence 1, Appli
C 31	14	37.8	1830121	4	US-10-329-960-1	Sequence 1, Appli
C 32	14	37.8	4403765	3	US-09-103-840A-2	Sequence 2, Appli
C 33	14	37.8	4411529	3	US-09-103-840A-1	Sequence 1, Appli
C 34	13	35.1	271	4	US-09-313-294A-695	Sequence 695, App
C 35	13	35.1	289	4	US-08-651-155B-12	Sequence 12, Appli
C 36	13	35.1	289	4	US-09-194-036B-12	Sequence 12, Appli
C 37	13	35.1	297	4	US-09-252-991A-8030	Sequence 8030, Ap
C 38	13	35.1	301	4	US-09-270-767-2113	Sequence 2113, Ap
C 39	13	35.1	301	4	US-09-270-767-17395	Sequence 17395, A
C 40	13	35.1	342	4	US-10-101-464A-147	Sequence 147, App
C 41	13	35.1	348	3	US-08-844-059-3	Sequence 3, Appli
C 42	13	35.1	348	3	US-09-431-202-3	Sequence 3, Appli
C 43	13	35.1	369	4	US-09-270-767-14734	Sequence 14734, A
C 44	13	35.1	410	4	US-09-221-017B-273	Sequence 273, App
C 45	13	35.1	423	4	US-09-252-991A-11388	Sequence 11388, A

ALIGNMENTS

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RESULT 1
US-08-864-357F-9
; Sequence 9, Application US/08864357F
; Patent No. 6255281
; GENERAL INFORMATION:
; APPLICANT: Clagen, Inc. & NIH
; TITLE OF INVENTION: Use of Recombinant Human Uteroglobulin in Treatment of Inflammato
; FILE REFERENCE: 116142/2
; CURRENT FILING DATE: 1997-05-28
; NUMBER OF SEQ ID NOS: 22
; SOFTWARE: PatentIn version 3.0
; SEQ ID NO 9
; LENGTH: 37
; TYPE: DNA
; ORGANISM: artificial
; FEATURE:
; OTHER INFORMATION: primer sequence
US-08-864-357F-9

Query Match      100.0%; Score 37; DB 3; Length 37;
Best Local Similarity 100.0%; Pred. No. 2.2e-12;
Matches 37; Conservative 0; Mismatches 0; Indels 0; Gaps 0;

QY
1 ATGAGAGAGATCGCTCAGTCTAGCTGTGCACTAAG 37
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US-08-864-357F-10/c
; Sequence 10, Application US/08864357F
; Patent No. 6255281
; GENERAL INFORMATION:
; APPLICANT: Clagen, Inc. & NIH
; TITLE OF INVENTION: Use of Recombinant Human Uteroglobulin in Treatment of Inflammato
; FILE REFERENCE: 116142/2
; CURRENT FILING DATE: 1997-05-28
; NUMBER OF SEQ ID NOS: 22
; SOFTWARE: PatentIn version 3.0
; SEQ ID NO 10
; LENGTH: 60
; TYPE: DNA
; ORGANISM: artificial
; FEATURE:
; OTHER INFORMATION: primer sequence
US-08-864-357F-10

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MOLECULE TYPE: CDNA
US-09-336-757-5

Query Match 37.8%; Score 14; DB 4; Length 283;
Best Local Similarity 100.0%; Pred. No. 43;
Matches 14; Conservative 0; Mismatches 0; Indels 0; Gaps 0;

QY 15 TCAGCTAGCCTGT 28
DB 269 TCAGCTAGCCTGT 256

RESULT 6
US-09-513-999C-3565/C
Sequence 3565 Application US/09513999C
Patent No. 6783961
GENERAL INFORMATION:
APPLICANT: Dumas Milne Edwards, J.B.
APPLICANT: Duclet, A.
APPLICANT: Giordano, J.Y.
TITLE OF INVENTION: Expressed Sequence Tags and Encoded Human Proteins.
Patent No. 6783961
FILE REFERENCE: 59 US2, REG
CURRENT APPLICATION NUMBER: US/09/513, 999C
CURRENT FILING DATE: 2000-02-24
PRIOR APPLICATION NUMBER: US 60/122,487
PRIOR FILING DATE: 1999-02-26
NUMBER OF SEQ ID NOS: 36681
SOFTWARE: Patent.pm
SEQ ID NO 3565
LENGTH: 341
TYPE: DNA
ORGANISM: Homo sapiens
FEATURE:
NAME/KEY: CDS
LOCATION: 22..339
US-09-513-999C-3565

Query Match 37.8%; Score 14; DB 4; Length 341;
Best Local Similarity 100.0%; Pred. No. 43;
Matches 14; Conservative 0; Mismatches 0; Indels 0; Gaps 0;

QY 16 CAGCTAGCCTGTG 29
DB 111 CAGCTAGCCTGTG 98

RESULT 7
US-09-621-976-12461
Sequence 12461 Application US/09621976
Patent No. 6639063
GENERAL INFORMATION:
APPLICANT: Dumas Milne Edwards, J.B.
APPLICANT: Gobert, S.
APPLICANT: Giordano, J.Y.
TITLE OF INVENTION: ESTs and Encoded Human Proteins.
FILE REFERENCE: GENSET, 054PR2
CURRENT APPLICATION NUMBER: US/09/621, 976
CURRENT FILING DATE: 2000-07-21
NUMBER OF SEQ ID NOS: 19335
SOFTWARE: Patent.pm
SEQ ID NO 12461
LENGTH: 422
TYPE: DNA
ORGANISM: Homo sapiens
US-09-621-976-12461

Query Match 37.8%; Score 14; DB 4; Length 422;
Best Local Similarity 100.0%; Pred. No. 43;
Matches 14; Conservative 0; Mismatches 0; Indels 0; Gaps 0;

QY 20 CTAGCCTGTGCAAC 33
CTAGCCTGTGCAAC 33

DB 354 CTAGCCTGTGCAAC 367

RESULT 8
US-08-463-115-26
Sequence 26 Application US/08463115
Patent No. 5703221
GENERAL INFORMATION:
APPLICANT: WILLIAM JOHN MARTIN
TITLE OF INVENTION: ISOLATED STEALTH VIRUSES
TITLE OF INVENTION: AND RELATED VACCINES
NUMBER OF SEQUENCES: 104
CORRESPONDENCE ADDRESS:
ADDRESS: Lyon & Lyon
STREET: 633 West Fifth Street
STREET: Suite 4700
CITY: Los Angeles
STATE: California
COUNTRY: U.S.A.
ZIP: 90071-2066
COMPUTER READABLE FORM:
MEDIUM TYPE: 3.5" Diskette, 1.44 Mb
MEDIUM TYPE: storage
COMPUTER: IBM Compatible
OPERATING SYSTEM: IBM P.C. DOS 5.0
SOFTWARE: FastSeq Version 1.5
CURRENT APPLICATION DATA:
APPLICATION NUMBER: US/08/463,115
FILING DATE: June 5, 1995
CLASSIFICATION: 435
PRIOR APPLICATION DATA:
PRIOR APPLICATION DATA: including application
PRIOR APPLICATION DATA: described below:
APPLICATION NUMBER: 08/157,811
FILING DATE: No. 5703221member 23, 1993
APPLICATION NUMBER: 07/887,502
FILING DATE: May 22, 1992
APPLICATION NUMBER: 07/704,814
FILING DATE: May 23, 1991
APPLICATION NUMBER: 07/763,039
FILING DATE: September 20, 1991
ATTORNEY/AGENT INFORMATION:
NAME: Warburg, Richard J.
REGISTRATION NUMBER: 32,327
REFERENCE/DOCKET NUMBER: 213/301
TELECOMMUNICATION INFORMATION:
TELEPHONE: (213) 489-1600
TELEFAX: (213) 955-0440
TELEX: 67-3510
INFORMATION FOR SEQ ID NO: 26:
SEQUENCE CHARACTERISTICS:
LENGTH: 553 base pairs
TYPE: nucleic acid
STRANDEDNESS: single
TOPOLOGY: linear
FEATURE:
OTHER INFORMATION:
US-08-463-115-26

Query Match 37.8%; Score 14; DB 1; Length 553;
Best Local Similarity 100.0%; Pred. No. 43;
Matches 14; Conservative 0; Mismatches 0; Indels 0; Gaps 0;

QY 19 TCTAGCCTGTGCAA 32
DB 310 TCTAGCCTGTGCAA 323

RESULT 9
US-08-465-388-26
Sequence 26 Application US/08465388
Patent No. 5753468
GENERAL INFORMATION:

APPLICANT: WILLIAM JOHN MARTIN
TITLE OF INVENTION: ISOLATED STEALTH VIRUSES
NUMBER OF SEQUENCES: 104
CORRESPONDENCE ADDRESSES:
ADDRESSEE: LYON & LYON
STREET: 633 West Fifth Street
CITY: Los Angeles
STATE: California
COUNTRY: U.S.A.
ZIP: 90071-2066
COMPUTER READABLE FORM:
MEDIUM TYPE: 3.5" Diskette, 1.44 Mb
MEDIUM TYPE: Storage
COMPUTER: IBM Compatible
OPERATING SYSTEM: IBM P.C. DOS 5.0
SOFTWARE: FastSeq Version 1.5
CURRENT APPLICATION DATA:
FILING DATE: June 5, 1995
CLASSIFICATION: 435
PRIOR APPLICATION DATA:
PRIOR APPLICATION DATA: including application
PRIOR APPLICATION DATA: described below:
APPLICATION NUMBER: 08/157,811
FILING DATE: No. 5753488ember 23, 1993
APPLICATION NUMBER: 07/887,502
FILING DATE: May 22, 1992
APPLICATION NUMBER: 07/704,814
FILING DATE: May 23, 1991
APPLICATION NUMBER: 07/763,039
FILING DATE: September 20, 1991
ATTORNEY/AGENT INFORMATION:
NAME: Warburg, Richard J.
REGISTRATION NUMBER: 32,327
REFERENCE/DOCKET NUMBER: 213/300
TELECOMMUNICATION INFORMATION:
TELEPHONE: (213) 489-1600
TELEFAX: (213) 955-0440
TELEX: 67-3510
INFORMATION FOR SEQ ID NO: 26:
SEQUENCE CHARACTERISTICS:
LENGTH: 553 base pairs
TYPE: nucleic acid
STRANDEDNESS: single
TOPOLOGY: linear
FEATURE:
OTHER INFORMATION:
US-08-463-388-26

Query Match 37.8%; Score 14; DB 1; Length 553;
Best Local Similarity 100.0%; Pred. No. 43;
Matches 14; Conservative 0; Mismatches 0; Indels 0; Gaps 0;

QY 19 TCTAGCCTGTGCA 32
DB 310 TCTAGCCTGTGCA 323

RESULT 10
US-08-463-115-18
Sequence 18, Application US/08463115
Patent No. 5703221
GENERAL INFORMATION:
APPLICANT: WILLIAM JOHN MARTIN
TITLE OF INVENTION: ISOLATED STEALTH VIRUSES
NUMBER OF SEQUENCES: 104
CORRESPONDENCE ADDRESSES:
ADDRESSEE: LYON & LYON
STREET: 633 West Fifth Street
CITY: Los Angeles
STATE: California
COUNTRY: U.S.A.
ZIP: 90071-2066
COMPUTER READABLE FORM:
MEDIUM TYPE: 3.5" Diskette, 1.44 Mb
MEDIUM TYPE: Storage
COMPUTER: IBM Compatible

CITY: Los Angeles
STATE: California
COUNTRY: U.S.A.
ZIP: 90071-2066
COMPUTER READABLE FORM:
MEDIUM TYPE: 3.5" Diskette, 1.44 Mb
MEDIUM TYPE: Storage
COMPUTER: IBM Compatible
OPERATING SYSTEM: IBM P.C. DOS 5.0
SOFTWARE: FastSeq Version 1.5
CURRENT APPLICATION DATA:
FILING DATE: June 5, 1995
CLASSIFICATION: 435
PRIOR APPLICATION DATA:
PRIOR APPLICATION DATA: including application
PRIOR APPLICATION DATA: described below:
APPLICATION NUMBER: 08/157,811
FILING DATE: No. 5703221ember 23, 1993
APPLICATION NUMBER: 07/887,502
FILING DATE: May 22, 1992
APPLICATION NUMBER: 07/704,814
FILING DATE: May 23, 1991
APPLICATION NUMBER: 07/763,039
FILING DATE: September 20, 1991
ATTORNEY/AGENT INFORMATION:
NAME: Warburg, Richard J.
REGISTRATION NUMBER: 32,327
REFERENCE/DOCKET NUMBER: 213/301
TELECOMMUNICATION INFORMATION:
TELEPHONE: (213) 489-1600
TELEFAX: (213) 955-0440
TELEX: 67-3510
INFORMATION FOR SEQ ID NO: 18:
SEQUENCE CHARACTERISTICS:
LENGTH: 715 base pairs
TYPE: nucleic acid
STRANDEDNESS: single
TOPOLOGY: linear
FEATURE:
OTHER INFORMATION:
US-08-463-115-18

Query Match 37.8%; Score 14; DB 1; Length 715;
Best Local Similarity 100.0%; Pred. No. 43;
Matches 14; Conservative 0; Mismatches 0; Indels 0; Gaps 0;

QY 19 TCTAGCCTGTGCA 32
DB 325 TCTAGCCTGTGCA 338

RESULT 11
US-08-463-388-18
Sequence 18, Application US/08463388
Patent No. 5753488
GENERAL INFORMATION:
APPLICANT: WILLIAM JOHN MARTIN
TITLE OF INVENTION: ISOLATED STEALTH VIRUSES
NUMBER OF SEQUENCES: 104
CORRESPONDENCE ADDRESSES:
ADDRESSEE: LYON & LYON
STREET: 633 West Fifth Street
CITY: Los Angeles
STATE: California
COUNTRY: U.S.A.
ZIP: 90071-2066
COMPUTER READABLE FORM:
MEDIUM TYPE: 3.5" Diskette, 1.44 Mb
MEDIUM TYPE: Storage
COMPUTER: IBM Compatible

OPERATING SYSTEM: IBM P.C. DOS 5.0
SOFTWARE: FastSeq Version 1.5
CURRENT APPLICATION DATA:
APPLICATION NUMBER: US/08/465,388
FILING DATE: June 5, 1995
CLASSIFICATION: 435
PRIOR APPLICATION DATA: including application
PRIOR APPLICATION DATA: described below:
APPLICATION NUMBER: 08/157,811
FILING DATE: No. 5753486 September 23, 1993
APPLICATION NUMBER: 07/887,502
FILING DATE: May 22, 1992
APPLICATION NUMBER: 07/704,814
FILING DATE: May 23, 1991
APPLICATION NUMBER: 07/763,039
FILING DATE: September 20, 1991
ATTORNEY/AGENT INFORMATION:
NAME: Warburg, Richard J.
REGISTRATION NUMBER: 32,327
REFERENCE/DOCKET NUMBER: 213/300
TELECOMMUNICATION INFORMATION:
TELEPHONE: (213) 489-1600
TELEFAX: (213) 955-0440
TELEX: 67-3510
INFORMATION FOR SEQ ID NO: 18:
SEQUENCE CHARACTERISTICS:
LENGTH: 715 base pairs
TYPE: nucleic acid
STRANDEDNESS: single
TOPOLOGY: linear
FEATURE:
OTHER INFORMATION:
US-08-465-388-18

Query Match 37.8%; Score 14; DB 1; Length 715;
Best Local Similarity 100.0%; Pred. No. 43;
Matches 14; Conservative 0; Mismatches 0; Indels 0; Gaps 0;

QY 19 TCTAGCCTGTGCA 32
DB 325 TCTAGCCTGTGCA 338

RESULT 12
US-09-252-991A-5999/c
Sequence 5999, Application US/09252991A
PATENT No. 6551795
GENERAL INFORMATION:
APPLICANT: Marc J. Rubenfield et al.
TITLE OF INVENTION: NUCLEIC ACID AND AMINO ACID SEQUENCES RELATING TO PSEUDOMONAS
FILE REFERENCE: 107196.116
CURRENT APPLICATION NUMBER: US/09/252,991A
PRIOR FILING DATE: 1999-02-18
PRIOR APPLICATION NUMBER: US 60/074,788
PRIOR FILING DATE: 1998-02-18
PRIOR APPLICATION NUMBER: US 60/094,190
PRIOR FILING DATE: 1998-07-27
NUMBER OF SEQ ID NOS: 33142
SEQ ID NO 5999
LENGTH: 1374
TYPE: DNA
ORGANISM: Pseudomonas aeruginosa
US-09-252-991A-5999

Query Match 37.8%; Score 14; DB 4; Length 1374;
Best Local Similarity 100.0%; Pred. No. 43;
Matches 14; Conservative 0; Mismatches 0; Indels 0; Gaps 0;

QY 4 GAGAAGATGCTCA 17
DB 78 GAGAAGATGCTCA 65

RESULT 13
US-08-463-115-2
Sequence 2, Application US/08463115
PATENT No. 5703221
GENERAL INFORMATION:
APPLICANT: WILLIAM JOHN MARTIN
TITLE OF INVENTION: ISOLATED STEALTH VIRUSES
NUMBER OF SEQUENCES: 104
CORRESPONDENCE ADDRESS:
ADDRESSER: Lyon & Lyon
STREET: 633 West Fifth Street
CITY: Los Angeles
STATE: California
COUNTRY: U.S.A.
ZIP: 90071-2066
COMPUTER READABLE FORM:
MEDIUM TYPE: 3.5" Diskette, 1.44 MB
MEDIUM TYPE: storage
COMPUTER: IBM Compatible
OPERATING SYSTEM: IBM P.C. DOS 5.0
SOFTWARE: FastSeq Version 1.5
CURRENT APPLICATION DATA:
APPLICATION NUMBER: US/08/463,115
FILING DATE: June 5, 1995
CLASSIFICATION: 435
PRIOR APPLICATION DATA: including application
PRIOR APPLICATION DATA: described below:
APPLICATION NUMBER: 08/157,811
FILING DATE: No. 5703221 September 23, 1993
APPLICATION NUMBER: 07/887,502
FILING DATE: May 22, 1992
APPLICATION NUMBER: 07/704,814
FILING DATE: May 23, 1991
APPLICATION NUMBER: 07/763,039
FILING DATE: September 20, 1991
ATTORNEY/AGENT INFORMATION:
NAME: Warburg, Richard J.
REGISTRATION NUMBER: 32,327
REFERENCE/DOCKET NUMBER: 213/301
TELECOMMUNICATION INFORMATION:
TELEPHONE: (213) 489-1600
TELEFAX: (213) 955-0440
TELEX: 67-3510
INFORMATION FOR SEQ ID NO: 2:
SEQUENCE CHARACTERISTICS:
LENGTH: 1540 base pairs
TYPE: nucleic acid
STRANDEDNESS: single
TOPOLOGY: linear
US-08-463-115-2

Query Match 37.8%; Score 14; DB 1; Length 1540;
Best Local Similarity 100.0%; Pred. No. 43;
Matches 14; Conservative 0; Mismatches 0; Indels 0; Gaps 0;

QY 19 TCTAGCCTGTGCA 32
DB 997 TCTAGCCTGTGCA 1010

RESULT 14
US-08-465-388-2
Sequence 2, Application US/08465388
PATENT No. 5753488
GENERAL INFORMATION:
APPLICANT: WILLIAM JOHN MARTIN
TITLE OF INVENTION: ISOLATED STEALTH VIRUSES
NUMBER OF SEQUENCES: AND RELATED VACCINES

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OM nucleic - nucleic search, using sw model

Run on: December 22, 2004, 23:36:53, Search time 518.67 Seconds
(without alignments)
397.214 Million cell updates/sec

Title: US-09-898-616A-4

Perfect score: 37
Sequence: 1 atgagaagatcgctcagctcagctcgtgaactaag 37

Scoring table: OLIGO_NUC
Gapop 60.0, Gapext 60.0

Searched: 4105333 seqs, 2784095677 residues

Word size: 0

Total number of hits satisfying chosen parameters: 8210666

Minimum DB seq length: 0
Maximum DB seq length: 2000000000

Post-processing: Listing first 45 summaries

Database: Published Applications NA:

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3: /cgn2_6/ptodata/1/pubna/US06_NEW_PUB.seq:*
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11: /cgn2_6/ptodata/1/pubna/US09C_PUBCOMB.seq:*
12: /cgn2_6/ptodata/1/pubna/US09C_PUBCOMB.seq:*
13: /cgn2_6/ptodata/1/pubna/US10_PUBCOMB.seq:*
14: /cgn2_6/ptodata/1/pubna/US10B_PUBCOMB.seq:*
15: /cgn2_6/ptodata/1/pubna/US10C_PUBCOMB.seq:*
16: /cgn2_6/ptodata/1/pubna/US10D_PUBCOMB.seq:*
17: /cgn2_6/ptodata/1/pubna/US10E_PUBCOMB.seq:*
18: /cgn2_6/ptodata/1/pubna/US10F_PUBCOMB.seq:*
19: /cgn2_6/ptodata/1/pubna/US11_NEW_PUB.seq:*
20: /cgn2_6/ptodata/1/pubna/US60_NEW_PUB.seq:*
21: /cgn2_6/ptodata/1/pubna/US60_PUBCOMB.seq:*

Pred. No. is the number of results predicted by chance to have a score greater than or equal to the score of the result being printed, and is derived by analysis of the total score distribution.

SUMMARIES

Result No.	Score	Query Match	Length	ID	Description
1	37	100.0	37	9	US-09-861-688-9
2	37	100.0	37	15	US-09-898-616A-4
3	37	100.0	37	15	US-10-187-498A-4
4	37	100.0	37	16	US-10-647-971-8
5	37	100.0	60	9	US-09-861-688-10
6	37	100.0	60	15	US-09-898-616A-5
7	37	100.0	60	15	US-10-187-498A-5
8	37	100.0	60	16	US-10-647-971-9
9	16	43.2	33706	13	US-10-087-192-1750
10	16	43.2	78268	13	US-10-087-192-742
11	15	40.5	261	18	US-10-674-124A-5867
12	15	40.5	389	16	US-10-424-599-75149

13	15	40.5	477	10	US-09-918-995-1312	Sequence 1312, Ap
14	15	40.5	574	13	US-10-027-632-131785	Sequence 131785, A
15	15	40.5	574	13	US-10-027-632-131785	Sequence 131785, A
16	15	40.5	574	13	US-10-027-632-131785	Sequence 131785, A
17	15	40.5	574	13	US-10-027-632-131785	Sequence 131785, A
18	15	40.5	679	16	US-10-424-599-82560	Sequence 82560, A
19	15	40.5	686	16	US-10-424-599-109965	Sequence 109965, A
20	15	40.5	727	18	US-10-425-115-90184	Sequence 90184, A
21	15	40.5	1043	16	US-09-397-945-82	Sequence 82, Ap1
22	15	40.5	1043	16	US-10-653-595-82	Sequence 82, Ap1
23	15	40.5	1089	15	US-10-195-730-20	Sequence 20, Ap1
24	15	40.5	1089	17	US-10-799-747-20	Sequence 20, Ap1
25	15	40.5	1224	17	US-10-437-963-94846	Sequence 94846, A
26	15	40.5	1436	17	US-10-437-963-46892	Sequence 46892, A
27	15	40.5	1946	17	US-10-767-701-14898	Sequence 14898, A
28	15	40.5	2201	10	US-09-791-254-4	Sequence 4, Ap1
29	15	40.5	2814	17	US-10-437-963-94851	Sequence 94851, A
30	15	40.5	2814	17	US-10-437-963-94806	Sequence 94806, A
31	15	40.5	2835	17	US-10-437-963-94776	Sequence 94776, A
32	15	40.5	2889	17	US-10-437-963-94807	Sequence 94807, A
33	15	40.5	2952	17	US-10-437-963-94766	Sequence 94766, A
34	15	40.5	3362	17	US-10-437-963-94707	Sequence 94707, A
35	15	40.5	3396	17	US-10-437-963-94769	Sequence 94769, A
36	15	40.5	3414	17	US-10-437-963-94802	Sequence 94802, A
37	15	40.5	3501	17	US-10-437-963-94774	Sequence 94774, A
38	15	40.5	3558	17	US-10-437-963-94702	Sequence 94702, A
39	15	40.5	3561	17	US-10-437-963-94836	Sequence 94836, A
40	15	40.5	3591	17	US-10-437-963-94843	Sequence 94843, A
41	15	40.5	3618	17	US-10-437-963-94841	Sequence 94841, A
42	15	40.5	3677	17	US-10-437-963-94855	Sequence 94855, A
43	15	40.5	3687	17	US-10-437-963-47011	Sequence 47011, A
44	15	40.5	3711	17	US-10-437-963-94812	Sequence 94812, A
45	15	40.5	4065	10	US-09-791-254-1	Sequence 1, Ap1

ALIGNMENTS

RESULT 1
US-09-861-688-9
Sequence 9, Application US/09861688
Patent No. US20020173460A1
GENERAL INFORMATION:
APPLICANT: Clargen, Inc. & NIH
TITLE OF INVENTION: Use of Recombinant Human Uteroglobulin in Treatment of
TITLE OF INVENTION: Fibrotic Conditions
FILE REFERENCE: 116142/2
CURRENT APPLICATION NUMBER: US/09/861,688
CURRENT FILING DATE: 2001-05-21
PRIOR APPLICATION NUMBER: 08/864,357
PRIOR FILING DATE: 1997-05-28
NUMBER OF SEQ ID NOS: 22
SOFTWARE: PatentIn version 3.0
SEQ ID NO 9
LENGTH: 37
TYPE: DNA
ORGANISM: Artificial Sequence
FEATURE:
OTHER INFORMATION: primer sequence
US-09-861-688-9

Query Match 100.0%, Score 37, DB 9, Length 37,
Best Local Similarity 100.0%, Pred. No. 1.3e-12,
Matches 37, Conservative 0, Mismatches 0, Indels 0, Gaps 0,

1 ATGAGAAGATCGCTCAGCTCAGCTGCTGCAACTAAG 37
1 ATGAGAAGATCGCTCAGCTCAGCTGCTGCAACTAAG 37

RESULT 2
US-09-898-616A-4

Sequence 4, Application US/09898616A
Publication No. US20030109429A1
GENERAL INFORMATION:
APPLICANT: Clargen Inc.
APPLICANT: Pilon, Apple L.
APPLICANT: Welch, Richard W.
TITLE OF INVENTION: Method for the Production of Purified rhug for the Treatment of
FILE REFERENCE: 116142/00170
CURRENT APPLICATION NUMBER: US/09/898,616A
CURRENT FILING DATE: 2002-10-15
PRIORITY FILING DATE: 1997-05-28
NUMBER OF SEQ ID NOS: 10
SOFTWARE: Patent in version 3.1
SEQ ID NO 4
LENGTH: 37
TYPE: DNA
ORGANISM: Artificial Sequence
FEATURE:
OTHER INFORMATION: Original amino acid sequence obtained from cDNA. Current submitte
FEATURE:
OTHER INFORMATION: d sequence maximized for expression in E. coli.
NAME/KEY: misc:feature
OTHER INFORMATION: Original amino acid sequence obtained from cDNA. Current submitte
OTHER INFORMATION: d sequence maximized for expression in E. coli.
US-09-898-616A-4
Query Match 100.0%; Score 37; DB 10; Length 37;
Best Local Similarity 100.0%; Pred. No. 1.3e-12;
Matches 37; Conservative 0; Mismatches 0; Indels 0; Gaps 0;
CY 1 ATGAGAAGATCGCTCAGCTTACGCTGTGCACTAG 37
DB 1 ATGAGAAGATCGCTCAGCTTACGCTGTGCACTAG 37
RESULT 3
US-10-187-498A-4
Sequence 4, Application US/10187498A
Publication No. US20030207795A1
GENERAL INFORMATION:
APPLICANT: Clargen Inc.
APPLICANT: Pilon, Apple L.
APPLICANT: Welch, Richard W.
TITLE OF INVENTION: Method for the Production of Purified rhug for the Treatment of
FILE REFERENCE: 116142/00260
CURRENT APPLICATION NUMBER: US/10/187,498A
CURRENT FILING DATE: 2001-07-02
PRIORITY FILING DATE: 1997-05-28
NUMBER OF SEQ ID NOS: 10
SOFTWARE: Patent in version 3.1
SEQ ID NO 4
LENGTH: 37
TYPE: DNA
ORGANISM: Artificial Sequence
FEATURE:
OTHER INFORMATION: Original amino acid sequence obtained from cDNA. Current submitte
OTHER INFORMATION: d sequence maximized for expression in E. coli.
NAME/KEY: misc:feature
OTHER INFORMATION: Original amino acid sequence obtained from cDNA. Current submitte
OTHER INFORMATION: d sequence maximized for expression in E. coli.
US-10-187-498A-4
Query Match 100.0%; Score 37; DB 15; Length 37;
Best Local Similarity 100.0%; Pred. No. 1.3e-12;
Matches 37; Conservative 0; Mismatches 0; Indels 0; Gaps 0;
CY 1 ATGAGAAGATCGCTCAGCTTACGCTGTGCACTAG 37

DB 1 ATGAGAAGATCGCTCAGCTTACGCTGTGCACTAG 37
RESULT 4
US-10-647-371-8
Sequence 8, Application US/10647371
Publication No. US20040047857A1
GENERAL INFORMATION:
APPLICANT: Clargen, Inc. & NIH
APPLICANT: Pilon, Apple L.
TITLE OF INVENTION: Use of Recombinant Human Uteroglobin in Treatment of Inflammatory
FILE REFERENCE: 116142-85
CURRENT APPLICATION NUMBER: US/10/647,371
CURRENT FILING DATE: 2003-08-25
PRIORITY FILING DATE: 09/549,926
NUMBER OF SEQ ID NOS: 12
SOFTWARE: Patent in version 3.2
SEQ ID NO 8
LENGTH: 37
TYPE: DNA
ORGANISM: Artificial
FEATURE:
OTHER INFORMATION: Description of Artificial Sequence: primer sequence
US-10-647-371-8
Query Match 100.0%; Score 37; DB 16; Length 37;
Best Local Similarity 100.0%; Pred. No. 1.3e-12;
Matches 37; Conservative 0; Mismatches 0; Indels 0; Gaps 0;
CY 1 ATGAGAAGATCGCTCAGCTTACGCTGTGCACTAG 37
DB 1 ATGAGAAGATCGCTCAGCTTACGCTGTGCACTAG 37
RESULT 5
US-09-861-688-10/c
Sequence 10, Application US/09861688
Patent No. US20020173460A1
GENERAL INFORMATION:
APPLICANT: Clargen, Inc. & NIH
APPLICANT: Pilon, Apple L.
TITLE OF INVENTION: Use of Recombinant Human Uteroglobin in Treatment of
FILE REFERENCE: 116142/2
CURRENT APPLICATION NUMBER: US/09/861,688
CURRENT FILING DATE: 2002-05-21
PRIORITY FILING DATE: 1997-05-28
NUMBER OF SEQ ID NOS: 22
SOFTWARE: Patent in version 3.0
SEQ ID NO 10
LENGTH: 60
TYPE: DNA
ORGANISM: Artificial Sequence
FEATURE:
OTHER INFORMATION: primer sequence
US-09-861-688-10
Query Match 100.0%; Score 37; DB 9; Length 60;
Best Local Similarity 100.0%; Pred. No. 1.3e-12;
Matches 37; Conservative 0; Mismatches 0; Indels 0; Gaps 0;
CY 1 ATGAGAAGATCGCTCAGCTTACGCTGTGCACTAG 37
DB 37 ATGAGAAGATCGCTCAGCTTACGCTGTGCACTAG 1
RESULT 6
US-09-898-616A-5/c
Sequence 5, Application US/09898616A
Publication No. US20030109429A1
GENERAL INFORMATION:

APPLICANT: Claragen Inc.
APPLICANT: Pilon, Richard L
TITLE OF INVENTION: Method for the Production of Purified rhug for the Treatment of
FILE REFERENCE: 116142/00170
CURRENT APPLICATION NUMBER: US/09/898,616A
CURRENT FILING DATE: 2002-10-15
PRIOR APPLICATION NUMBER: US 08/864,357
PRIOR FILING DATE: 1997-05-28
NUMBER OF SEQ ID NOS: 10
SOFTWARE: PatentIn version 3.1
SEQ ID NO 5
LENGTH: 60
TYPE: DNA
ORGANISM: Artificial Sequence
FEATURE:
OTHER INFORMATION: Original amino acid sequence obtained from cDNA. Current submitte
FEATURE:
OTHER INFORMATION: d sequence maximized for expression in E. coli.
NAME/KEY: misc.feature
OTHER INFORMATION: Original amino acid sequence obtained from cDNA. Current submitte
OTHER INFORMATION: d sequence maximized for expression in E. coli.
US-09-898-616A-5

Query Match 100.0%; Score 37; DB 10; Length 60;
Best Local Similarity 100.0%; Pred. No. 1.3e-12;
Matches 37; Conservative 0; Mismatches 0; Indels 0; Gaps 0;

QY 1 ATGAGAGAGATCGCTCAGCTAGCCTGTGCAACTAG 37
DB 37 ATGAGAGAGATCGCTCAGCTAGCCTGTGCAACTAG 1

RESULT 7
US-10-187-498A-5/c
Sequence 5, Application US/10/87498A
Publication No. US2003020795A1
GENERAL INFORMATION:
APPLICANT: Claragen Inc.
APPLICANT: Pilon, Richard L
APPLICANT: Welch, Richard W
TITLE OF INVENTION: Method for the Production of Purified rhug for the Treatment of
FILE REFERENCE: 116142/00260
CURRENT APPLICATION NUMBER: US/10/187,498A
CURRENT FILING DATE: 2001-07-02
PRIOR APPLICATION NUMBER: US 08/864,357
PRIOR FILING DATE: 1997-05-28
NUMBER OF SEQ ID NOS: 10
SOFTWARE: PatentIn version 3.1
SEQ ID NO 5
LENGTH: 60
TYPE: DNA
ORGANISM: Artificial Sequence
FEATURE:
OTHER INFORMATION: Original amino acid sequence obtained from cDNA. Current submitte
OTHER INFORMATION: d sequence maximized for expression in E. coli.
NAME/KEY: misc.feature
OTHER INFORMATION: Original amino acid sequence obtained from cDNA. Current submitte
OTHER INFORMATION: d sequence maximized for expression in E. coli.
US-10-187-498A-5

Query Match 100.0%; Score 37; DB 15; Length 60;
Best Local Similarity 100.0%; Pred. No. 1.3e-12;
Matches 37; Conservative 0; Mismatches 0; Indels 0; Gaps 0;

QY 1 ATGAGAGAGATCGCTCAGCTAGCCTGTGCAACTAG 37
DB 37 ATGAGAGAGATCGCTCAGCTAGCCTGTGCAACTAG 1

RESULT 8
US-10-647-371-9/c
Sequence 9, Application US/10647371
Publication No. US20040047857A1
GENERAL INFORMATION:
APPLICANT: Claragen, Inc. & NIH
TITLE OF INVENTION: Use of Recombinant Human Uteroglobin in Treatment of Inflammatory
FILE REFERENCE: 116142-85
CURRENT APPLICATION NUMBER: US/10/647,371
CURRENT FILING DATE: 2003-08-25
PRIOR APPLICATION NUMBER: 09/549,926
PRIOR FILING DATE: 2000-04-14
NUMBER OF SEQ ID NOS: 12
SOFTWARE: PatentIn version 3.2
SEQ ID NO 9
LENGTH: 60
TYPE: DNA
ORGANISM: Artificial
FEATURE:
OTHER INFORMATION: Description of Artificial Sequence: primer sequence
US-10-647-371-9

Query Match 100.0%; Score 37; DB 16; Length 60;
Best Local Similarity 100.0%; Pred. No. 1.3e-12;
Matches 37; Conservative 0; Mismatches 0; Indels 0; Gaps 0;

QY 1 ATGAGAGAGATCGCTCAGCTAGCCTGTGCAACTAG 37
DB 37 ATGAGAGAGATCGCTCAGCTAGCCTGTGCAACTAG 1

RESULT 9
US-10-087-192-1750
Sequence 1750, Application US/10087192
Publication No. US20020182586A1
GENERAL INFORMATION:
APPLICANT: Morris, David W.
APPLICANT: Engelhard, Eric K.
TITLE OF INVENTION: NOVEL COMPOSITIONS AND METHODS FOR
FILE REFERENCE: 528452000122
CURRENT APPLICATION NUMBER: US/10/087,192
CURRENT FILING DATE: 2002-03-01
PRIOR APPLICATION NUMBER: US 09/747,377
PRIOR FILING DATE: 2000-12-22
PRIOR APPLICATION NUMBER: US 09/798,586
PRIOR FILING DATE: 2001-03-02
NUMBER OF SEQ ID NOS: 2059
SOFTWARE: FastSeq for Windows Version 4.0
SEQ ID NO 1750
LENGTH: 32706
TYPE: DNA
ORGANISM: Homo sapiens
US-10-087-192-1750

Query Match 43.2%; Score 16; DB 13; Length 32706;
Best Local Similarity 100.0%; Pred. No. 7.4; Indels 0; Gaps 0;
Matches 16; Conservative 0; Mismatches 0;

QY 19 TCTAGCCTGTGCAACT 34
DB 7293 TCTAGCCTGTGCAACT 7308

RESULT 10
US-10-087-192-742/c
Sequence 742, Application US/10087192
Publication No. US20020182586A1
GENERAL INFORMATION:
APPLICANT: Morris, David W.
APPLICANT: Engelhard, Eric K.
TITLE OF INVENTION: NOVEL COMPOSITIONS AND METHODS FOR

TITLE OF INVENTION: CANCER
FILE REFERENCE: 529452000122
CURRENT APPLICATION NUMBER: US/10/087,192
CURRENT FILING DATE: 2002-03-01
PRIOR APPLICATION NUMBER: US 09/747,377
PRIOR FILING DATE: 2000-12-22
PRIOR APPLICATION NUMBER: US 09/798,586
PRIOR FILING DATE: 2001-03-02
NUMBER OF SEQ ID NOS: 2059
SOFTWARE: FastSeq for Windows Version 4.0
SEQ ID NO 742
LENGTH: 78268
TYPE: DNA
ORGANISM: Homo sapiens
US-10-087-192-742

Query Match 43.2% Score 16; DB 13; Length 78268;
Best Local Similarity 100.0%; Pred. No. 7.2;
Matches 16; Conservative 0; Mismatches 0; Indels 0; Gaps 0;

QY 13 GCTGAGCTGTGCTGT 28
DB 29223 GCTGAGCTGTGCTGT 29208

RESULT 11
US-10-674-124A-5867/C
Sequence 5867, Application US/10674124A
Publication No. US2004019797A1
GENERAL INFORMATION:
APPLICANT: INOKO, Hidetoshi
APPLICANT: TAMURA, Gen
TITLE OF INVENTION: GENE MAPPING METHOD USING MICROSATELLITE.
TITLE OF INVENTION: GENETIC POLYMORPHISM MARKERS
FILE REFERENCE: ORIN-003CIP
CURRENT APPLICATION NUMBER: US/10/674,124A
CURRENT FILING DATE: 2003-09-26
PRIOR APPLICATION NUMBER: 10/257,511
PRIOR FILING DATE: 2003-03-07
PRIOR APPLICATION NUMBER: PCT/JP00/07621
PRIOR FILING DATE: 2000-10-30
PRIOR APPLICATION NUMBER: JP2000-112699
PRIOR FILING DATE: 2000-04-13
PRIOR APPLICATION NUMBER: JP2002-327516
PRIOR FILING DATE: 2002-09-28
PRIOR APPLICATION NUMBER: JP2002-383869
PRIOR FILING DATE: 2002-12-09
NUMBER OF SEQ ID NOS: 27110
SEQ ID NO 5867
LENGTH: 261
TYPE: DNA
ORGANISM: Homo sapiens
FEATURE:
OTHER INFORMATION: AC072028.5_149909
FEATURE:
OTHER INFORMATION: Located on chromosome 3
FEATURE:
OTHER INFORMATION: Distance between a terminus base of telomere on
OTHER INFORMATION: chromosomal short arm and 5'-terminus of this base
FEATURE:
OTHER INFORMATION: sequence: 147898939
FEATURE:
OTHER INFORMATION: Distance between 3'-terminus of neighbour sequence of
OTHER INFORMATION: sequence listing upward to telomere on chromosomal short arm and
OTHER INFORMATION: 5'-terminus of this base sequence: 130675
US-10-674-124A-5867

Query Match 40.5% Score 15; DB 18; Length 261;
Best Local Similarity 100.0%; Pred. No. 36;
Matches 15; Conservative 0; Mismatches 0; Indels 0; Gaps 0;

QY 19 TTAGCCTGTGCAAC 33
DB 240 TTAGCCTGTGCAAC 226

RESULT 12
US-10-424-599-75149/C
Sequence 75149, Application US/10424599
Publication No. US20040031072A1
GENERAL INFORMATION:
APPLICANT: La Rosa Thomas J
APPLICANT: Kovalic David K
APPLICANT: Zhou Yihua
APPLICANT: Cao Yongwei
TITLE OF INVENTION: Soy Nucleic Acid Molecules and Other Molecules Associated With
TITLE OF INVENTION: Plants and Uses Thereof for Plant Improvement
FILE REFERENCE: 38-21(53223)B
CURRENT APPLICATION NUMBER: US/10/424,599
CURRENT FILING DATE: 2003-04-28
NUMBER OF SEQ ID NOS: 285684
SEQ ID NO 75149
LENGTH: 389
TYPE: DNA
ORGANISM: Glycine max
FEATURE:
OTHER INFORMATION: Clone ID: PAT_MRT3847_38873C.1
US-10-424-599-75149

Query Match 40.5% Score 15; DB 16; Length 389;
Best Local Similarity 100.0%; Pred. No. 35;
Matches 15; Conservative 0; Mismatches 0; Indels 0; Gaps 0;

QY 15 TAGGTCTAGCCTGTG 29
DB 40 TAGGTCTAGCCTGTG 26

RESULT 13
US-09-918-995-1312
Sequence 1312, Application US/09918995
Publication No. US20030073623A1
GENERAL INFORMATION:
APPLICANT: Hyseq, Inc.
TITLE OF INVENTION: NOVEL NUCLEIC ACID SEQUENCES OBTAINED
TITLE OF INVENTION: FROM VARIOUS CDNA LIBRARIES
FILE REFERENCE: 20411-756
CURRENT APPLICATION NUMBER: US/09/918,995
CURRENT FILING DATE: 2001-07-30
PRIOR APPLICATION NUMBER: US/09/235,076
PRIOR FILING DATE: 1999-01-20
NUMBER OF SEQ ID NOS: 38054
SOFTWARE: FastSeq for Windows Version 3.0
SEQ ID NO 1312
LENGTH: 477
TYPE: DNA
ORGANISM: Homo sapiens
FEATURE:
NAME/KEY: misc_feature
LOCATION: (1)...(477)
OTHER INFORMATION: n = A,T,C or G
US-09-918-995-1312

Query Match 40.5% Score 15; DB 10; Length 477;
Best Local Similarity 100.0%; Pred. No. 35;
Matches 15; Conservative 0; Mismatches 0; Indels 0; Gaps 0;

QY 2 TGAGAGAGTCGCTC 16
DB 305 TGAGAGAGTCGCTC 319

RESULT 14
US-10-027-632-131785/C
Sequence 131785, Application US/10027632
Publication No. US20020198371A1
GENERAL INFORMATION:


```

APPLICANT: Mang, David G.
TITLE OF INVENTION: Identification and Mapping of Single Nucleotide
FILE REFERENCE: 108627.129
CURRENT APPLICATION NUMBER: US/10/027,632
PRIOR FILING DATE: 2002-04-30
PRIOR APPLICATION NUMBER: US 60/218,006
PRIOR FILING DATE: 2000-07-12
PRIOR APPLICATION NUMBER: US 60/198,676
PRIOR FILING DATE: 2000-04-20
PRIOR APPLICATION NUMBER: US 60/193,483
PRIOR FILING DATE: 2000-03-29
PRIOR APPLICATION NUMBER: US 60/185,218
PRIOR FILING DATE: 2000-02-24
PRIOR APPLICATION NUMBER: US 60/167,363
PRIOR FILING DATE: 1999-11-23
PRIOR APPLICATION NUMBER: US 60/156,358
PRIOR FILING DATE: 1999-09-28
PRIOR APPLICATION NUMBER: US 60/146,002
PRIOR FILING DATE: 1999-08-09
NUMBER OF SEQ ID NOS: 325720
SOFTWARE: FastSeq for Windows Version 4.0
SEQ ID NO 131785
LENGTH: 574
TYPE: DNA
ORGANISM: Human
US-10-027-632-131785

```

```

Query Match          40.5%; Score 15; DB 13; Length 574;
Best Local Similarity 100.0%; Pred.No. 35;
Matches 15; Conservative 0; Mismatches 0; Indels 0; Gaps 0;

```

```

QY      18 GCTAGCCTGTGCAA 32
         |||||
DB      40 GCTAGCCTGTGCAA 26

```

```

RESULT 15
US-10-027-632-131786/c
Sequence 131786, Application US/10027632
Publication No. US20020198371A1
GENERAL INFORMATION:
APPLICANT: Mang, David G.
TITLE OF INVENTION: Identification and Mapping of Single Nucleotide
FILE REFERENCE: 108627.129
CURRENT APPLICATION NUMBER: US/10/027,632
CURRENT FILING DATE: 2002-04-30
PRIOR APPLICATION NUMBER: US 60/218,006
PRIOR FILING DATE: 2000-07-12
PRIOR APPLICATION NUMBER: US 60/198,676
PRIOR FILING DATE: 2000-04-20
PRIOR APPLICATION NUMBER: US 60/193,483
PRIOR FILING DATE: 2000-03-29
PRIOR APPLICATION NUMBER: US 60/185,218
PRIOR FILING DATE: 2000-02-24
PRIOR APPLICATION NUMBER: US 60/167,363
PRIOR FILING DATE: 1999-11-23
PRIOR APPLICATION NUMBER: US 60/156,358
PRIOR FILING DATE: 1999-09-28
PRIOR APPLICATION NUMBER: US 60/146,002
PRIOR FILING DATE: 1999-08-09
NUMBER OF SEQ ID NOS: 325720
SOFTWARE: FastSeq for Windows Version 4.0
SEQ ID NO 131786
LENGTH: 574
TYPE: DNA
ORGANISM: Human
US-10-027-632-131786

```

```

Query Match          40.5%; Score 15; DB 13; Length 574;
Best Local Similarity 100.0%; Pred.No. 35;
Matches 15; Conservative 0; Mismatches 0; Indels 0; Gaps 0;

```

```

QY      18 GCTAGCCTGTGCAA 32
         |||||
DB      40 GCTAGCCTGTGCAA 26

```

```

Search completed: December 23, 2004, 05:19:30
Job time : 520.67 secs

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